

116809

U.S. DEPARTMENT OF COMMERCE
Patent and Trademark Office

SEARCH REQUEST FORM

Requestor's

Name: Hong Liu

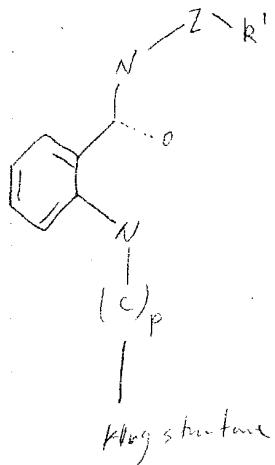
Serial

Number: 09/831,506Date: 3/11/04Phone: 2-0669Art Unit: 1624Ren 5c11

Search Topic:

Please write a detailed statement of search topic. Describe specifically as possible the subject matter to be searched. Define any terms that may have a special meaning. Give examples or relevant citations, authors, keywords, etc., if known. For sequences, please attach a copy of the sequence. You may include a copy of the broadest and/or most relevant claim(s).

Barb please?



anthranilic acid amides

A Huth D Seidelmann K Thierauch

STAFF USE ONLY

Date completed: 3-16-04Searcher: ACBTerminal time: 31Elapsed time: prep 30CPU time: 1Total time: Number of Searches: Number of Databases:

Search Site

 STIC

Vendors

 IG CM-1 522 STN Pre-S Dialog

Type of Search

 N.A. Sequence Geninfo A.A. Sequence SDC 3 Structure DARC/Questel Bibliographic Other

=> fil reg; d stat que 127; fil cap1;d que nos 128; fil uspatf; d que nos 129
FILE 'REGISTRY' ENTERED AT 15:17:55 ON 16 MAR 2004
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 15 MAR 2004 HIGHEST RN 663595-21-9
DICTIONARY FILE UPDATES: 15 MAR 2004 HIGHEST RN 663595-21-9

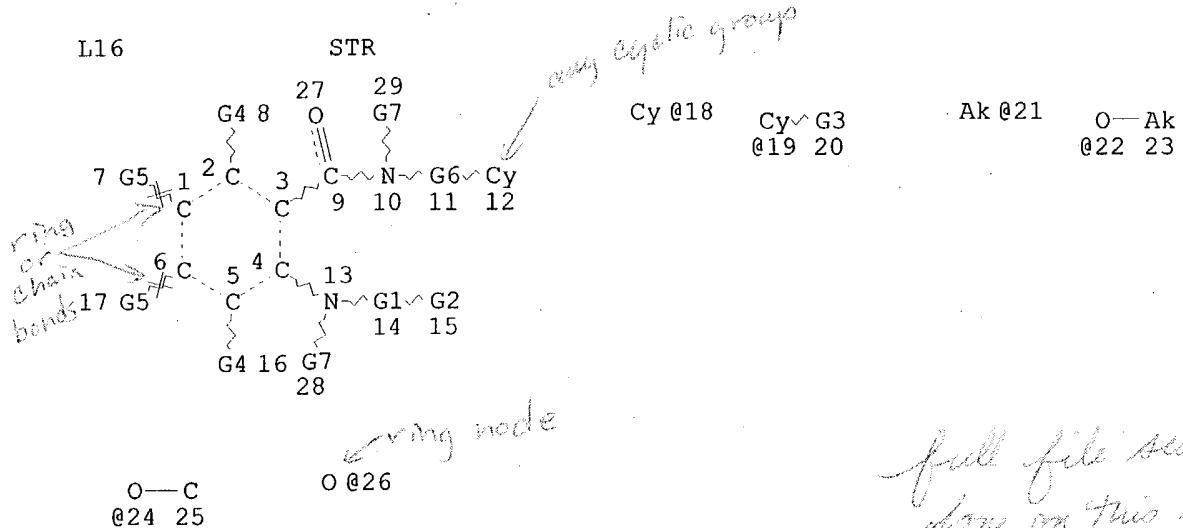
TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:

<http://www.cas.org/ONLINE/DBSS/registryss.html>



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REP G1=(1-4) C
VAR G2=18/19
VAR G3=X/21/22/0H
VAR G4=H/X/24/C
VAR G5=H/X/24/C/26
REP G6=(0-9) C
VAR G7=H/21
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VAR G:=H/21

NODE ATTRIBUTES

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CONNECT IS E1 PC

CONNECT IS E1 RC
CONNECT IS E1 RC

CONNECT IS E1

CONNECT IS
READY TO USE

DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

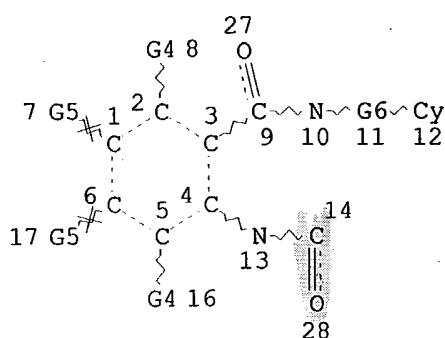
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NUMBER OF NODES IS 29

full file search
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with structure on next
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of answer set

STEREO ATTRIBUTES: NONE

L18 STR



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 @24 25

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REP G6=(0-9) C

NODE ATTRIBUTES:

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DEFAULT MLEVEL IS ATOM

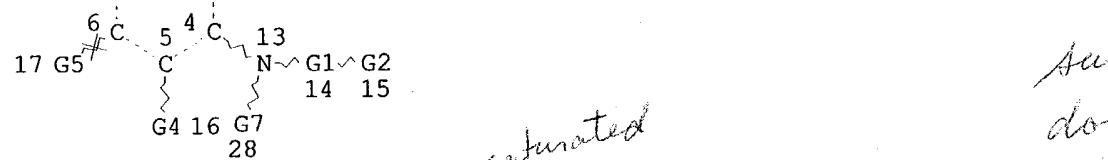
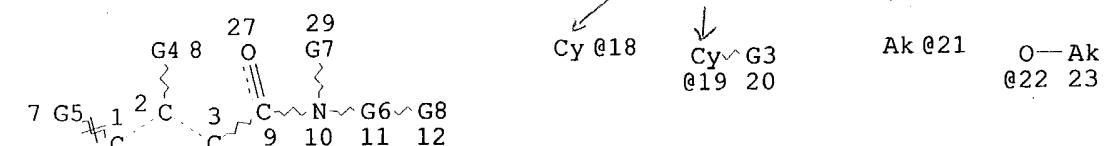
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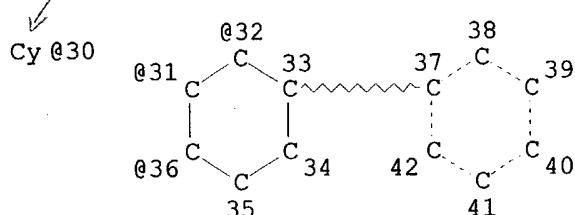
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NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L21 425 SEA FILE=REGISTRY SSS FUL L16 NOT L18
 L25 STR

O—C O @26
 @24 25



subset search
 done on this
 structure

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VAR G2=18/19

VAR G3=X/21/22/OH

VAR G4=H/X/24/C

VAR G5=H/X/24/C/26

REP G6=(0-9) C
VAR G7=H/21
VAR G8=30/32/31/36

NODE ATTRIBUTES:

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CONNECT IS E1 RC AT 18
CONNECT IS E1 RC AT 21
CONNECT IS E1 RC AT 23
DEFAULT MLEVEL IS ATOM
GGCAT IS UNS AT 18
GGCAT IS UNS AT 19
GGCAT IS UNS AT 30
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 42

STEREO ATTRIBUTES: NONE

L27 325 SEA FILE=REGISTRY SUB=L21 SSS FUL L25

100.0% PROCESSED 425 ITERATIONS
SEARCH TIME: 00.00.01

325 ANSWERS

FILE 'CAPLUS' ENTERED AT 15:17:55 ON 16 MAR 2004
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FILE COVERS 1907 - 16 Mar 2004 VOL 140 ISS 12
FILE LAST UPDATED: 15 Mar 2004 (20040315/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

L16	STR
L18	STR
L21	425 SEA FILE=REGISTRY SSS FUL L16 NOT L18
L25	STR
L27	325 SEA FILE=REGISTRY SUB=L21 SSS FUL L25
L28	43 SEA FILE=CAPLUS ABB=ON L27

FILE 'USPATFULL' ENTERED AT 15:17:55 ON 16 MAR 2004
CA INDEXING COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

FILE COVERS 1971 TO PATENT PUBLICATION DATE: 16 Mar 2004 (20040316/PD)
FILE LAST UPDATED: 16 Mar 2004 (20040316/ED)
HIGHEST GRANTED PATENT NUMBER: US6708338
HIGHEST APPLICATION PUBLICATION NUMBER: US2004049824
CA INDEXING IS CURRENT THROUGH 16 Mar 2004 (20040316/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 16 Mar 2004 (20040316/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2004
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2004

>>> USPAT2 is now available. USPATFULL contains full text of the <<<
>>> original, i.e., the earliest published granted patents or <<<
>>> applications. USPAT2 contains full text of the latest US <<<
>>> publications, starting in 2001, for the inventions covered in <<<
>>> USPATFULL. A USPATFULL record contains not only the original <<<
>>> published document but also a list of any subsequent <<<
>>> publications. The publication number, patent kind code, and <<<
>>> publication date for all the US publications for an invention <<<
>>> are displayed in the PI (Patent Information) field of USPATFULL <<<
>>> records and may be searched in standard search fields, e.g., /PN, <<<
>>> /PK, etc. <<<

>>> USPATFULL and USPAT2 can be accessed and searched together <<<
>>> through the new cluster USPATALL. Type FILE USPATALL to <<<
>>> enter this cluster. <<<
>>> <<<
>>> Use USPATALL when searching terms such as patent assignees, <<<
>>> classifications, or claims, that may potentially change from <<<
>>> the earliest to the latest publication. <<<

This file contains CAS Registry Numbers for easy and accurate substance identification.

L16 STR
L18 STR
L21 425 SEA FILE=REGISTRY SSS FUL L16 NOT L18
L25 STR
L27 325 SEA FILE=REGISTRY SUB=L21 SSS FUL L25
L29 14 SEA FILE=USPATFULL ABB=ON L27

=> dup rem 128,129
FILE 'CAPLUS' ENTERED AT 15:17:59 ON 16 MAR 2004
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FILE 'USPATFULL' ENTERED AT 15:17:59 ON 16 MAR 2004
CA INDEXING COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)
PROCESSING COMPLETED FOR L28
PROCESSING COMPLETED FOR L29
L31 55 DUP REM L28 L29 (2 DUPLICATES REMOVED)
ANSWERS '1-43' FROM FILE CAPLUS
ANSWERS '44-55' FROM FILE USPATFULL

~~d ibib ed abs hitstr 1-55; fil cao; d que nos 130~~

L31 ANSWER 1 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 1
ACCESSION NUMBER: 2003:950057 CAPLUS
DOCUMENT NUMBER: 140:16647
TITLE: Preparation of 2-aminopyridine-3-carboxamides as
remedies for angiogenesis mediated diseases

INVENTOR(S): Askew, Benny; Adams, Jeffrey; Book, Shon; Chen, Guoqing; Dipietro, Lucian V.; Elbaum, Daniel; Germain, Julie; Geuns-Meyer, Stephanie D.; Habgood, Gregory J.; Handley, Michael; Huang, Qi; Kim, Tae-seong; Li, Aiwen; Nishimura, Nobuko; Nomak, Rana; Patel, Vinod F.; Riahi, Babak; Kim, Joseph L.; Xi, Ning; Yang, Kevin; Yuan, Chester Chenguang

PATENT ASSIGNEE(S): Amgen Inc., USA

SOURCE: U.S. Pat. Appl. Publ., 252 pp., Cont.-in-part of U.S. Ser. No. 46,681.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

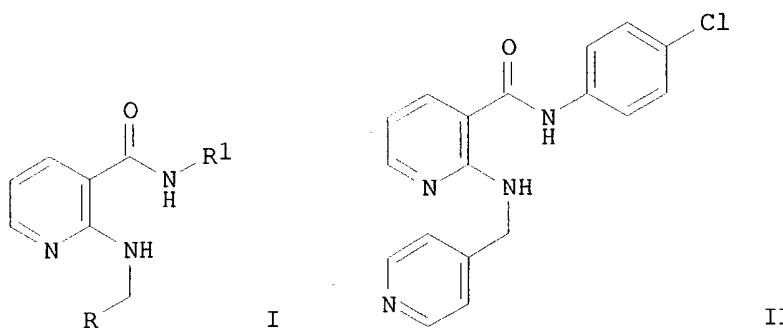
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003225106	A1	20031204	US 2002-197974	20020717
US 2003125339	A1	20030703	US 2002-46681	20020110
WO 2004007458	A1	20040122	WO 2003-US22417	20030715
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			US 2001-261339P	P 20010112
			US 2001-323764P	P 20010919
			US 2002-46681	A2 20020110
			US 2002-197974	A 20020717

OTHER SOURCE(S): MARPAT 140:16647

ED Entered STN: 05 Dec 2003
GI



AB The title compds. [I; R = (un)substituted 4-pyridyl, 2-pyridyl, 4-pyrimidinyl, 4-quinolyl, etc.; R1 = (un)substituted aryl, cycloalkyl, 5-6 membered heteroaryl, 9-10 membered bicyclic and 11-14 membered tricyclic heterocycl], which are effective for prophylaxis and treatment of diseases and other maladies or conditions involving, cancer and the

like, were prepd. Thus, the title compd. II was prepd. from 2-aminonicotinic acid, 4-chloroaniline, and 4-pyridinecarboxaldehyde. The compds. I showed inhibition of KDR kinase at < 50 μ M. Many compds. I inhibited VEGF-stimulated HUVEC proliferation at a level below 50 nM. Pharmaceutical compn. comprising the compd. I is claimed.

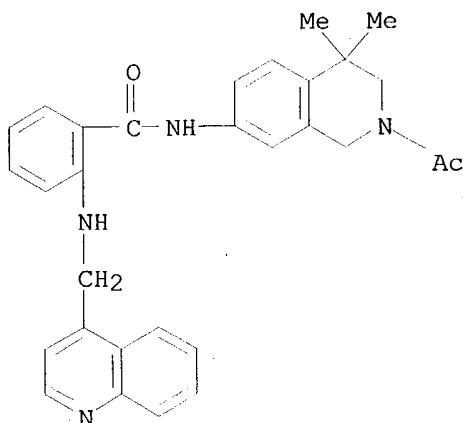
IT 453564-10-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of 2-aminopyridine-3-carboxamides for treating angiogenesis mediated diseases)

RN 453564-10-8 CAPLUS

CN Benzamide, N-(2-acetyl-1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)



L31 ANSWER 2 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 2

ACCESSION NUMBER: 2000:769086 CAPLUS

DOCUMENT NUMBER: 133:335159

TITLE: Preparation of N-pyridinyl-2-[(thienylcarbonyl)amino]benzamides and analogs as anticoagulants

INVENTOR(S): Arnaiz, Damian O.; Chou, Yuo-ling; Griedel, Brian D.; Karanjawala, Rushad E.; Kochanny, Monica J.; Lee, Wheeseong; Liang, Amy Mei; Morrissey, Michael M.; Phillips, Gary B.; Sacchi, Karna Lyn; Sakata, Steven T.; Shaw, Kenneth J.; Snider, R. Michael; Wu, Shung C.; Ye, Bin; Zhao, Zuchun

PATENT ASSIGNEE(S): Berlex Laboratories, Inc., USA

SOURCE: U.S., 113 pp., Cont.-in-part of U.S. Ser. No. 994,284, abandoned.

CODEN: USXXAM

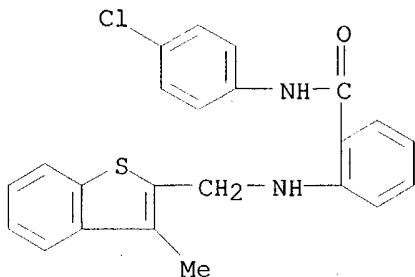
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6140351	A	20001031	US 1998-187459	19981105
CA 2315070	AA	19990701	CA 1998-2315070	19981127
WO 9932477	A1	19990701	WO 1998-EP7650	19981127
W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,			



REFERENCE COUNT: 45 THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

~~LS1~~ ANSWER 3 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:143094 CAPLUS

TITLE: Preparation of substituted (2S)-(arylamino)-3-(biphenyl-4-yl)propionic acids as antagonists of factor IX for inhibiting the intrinsic pathway of blood coagulation

INVENTOR(S): Mjalli, Adnan M. M.; Andrews, Robert C.; Guo, Xiao-chuan; Christen, Daniel Peter; Gohimmukkula, Devi Reddy; Huang, Guoxiang; Rothlein, Robert; Tyagi, Sameer; Yaramasu, Tripura; Behme, Christopher

PATENT ASSIGNEE(S): Transtech Pharma, Inc., USA

SOURCE: PCT Int. Appl., 326 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004014844	A2	20040219	WO 2003-US25045	20030808
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: US 2002-402272P P 20020809

ED Entered STN: 22 Feb 2004

AB The title compds. Ar2XCH(VAr1)(CH2)cG [I; c = 0-2; G = H, CO2R1, CH2OR1, COR1, CR1:NOR2, an acid isostere (wherein R1, R2 = H, alkyl, aryl, etc.); V = (CH2)bO(CH2)a, (CH2)bNR7(CH2)a, (CH2)bO, (CH2)bNR7, (CH2)a, a bond (a = 0-2; b = 1-2; R7 = H, alkyl, aryl, etc.); X = NR8, COR8, NR8CO, etc. (R8 = H, alkyl, aryl, etc.); Ar1 = (un)substituted aryl, heteroaryl, cycloalkylaryl, etc.; Ar2 = (un)substituted aryl or heteroaryl], useful as antagonists, or more preferably, partial antagonists of factor IX and thus, may be used to inhibit the intrinsic pathway of blood coagulation, were prep'd. Thus, reacting Me 2-L-amino-3-biphenyl-4-yl-propionate with isoquinoline-3-carboxylic acid followed by hydrolysis afforded 81% 3-biphenyl-4-yl-(2S)-[(isoquinoline-3-carbonyl)amino]propionic acid. The compds. I inhibit factor IX with IC50 of less than 30 .mu.M, and are useful in a variety of applications including the management, treatment

NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
 UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
 FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
 CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

AU 9918759 A1 19990712 AU 1999-18759 19981127

AU 751856 B2 20020829

EP 1040108 A1 20001004

EP 1998-963519 19981127

EP 1040108 B1 20040225

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, FI

JP 2001526283 T2 20011218 JP 2000-525414 19981127

NZ 503809 A 20020426 NZ 1998-503809 19981127

ZA 9811599 A 19990817 ZA 1998-11599 19981217

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US 6380221 B1 20020430 US 2000-631450 20000803

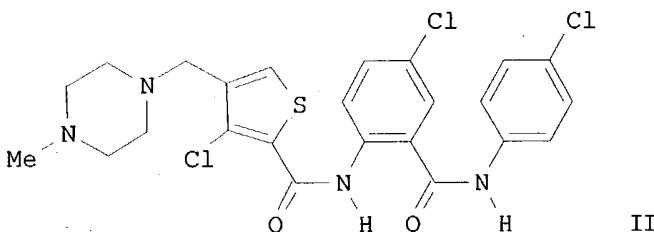
US 6498185 B1 20021224 US 2000-631452 20000803

PRIORITY APPLN. INFO.: US 1997-994284 B2 19971219
 US 1998-187459 A 19981105
 WO 1998-EP7650 W 19981127

OTHER SOURCE(S): MARPAT 133:335159

ED Entered STN: 02 Nov 2000

GI



AB RÉZDR3 [I; D, E = Z1NR5C(:X), Z1NR5SO0-2, etc.; R, R3 = (un)substituted heterocycll or -aryl; R5 = H, (ar)alkyl, aryl; X = O, S, H2; Z = (un)substituted heterocyclene or -arylene; Z1 = bond, alkylene, alkylidene, etc.] were prep'd. as factor Xa, thrombin, and prothrombinase inhibitors. Thus, H2NZCONHC6H4Cl-4 (Z = 4-chloro-1,2-phenylene) (prepn. given) was N-acylated by 3-chloro-4-chloromethyl-2-thiophenecarbonyl chloride and the product aminated by 1-methylpiperazine to give title compd. II. Data for biol. activity of I were given.

IT 229339-81-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of N-pyridinyl-2-[(thienylcarbonyl)amino]benzamides and analogs as anticoagulants)

RN 229339-81-5 CAPLUS

CN Benzamide, N-(4-chlorophenyl)-2-[(3-methylbenzo[b]thien-2-yl)methyl]amino]- (9CI) (CA INDEX NAME)

and/or control of diseases caused in part by the intrinsic clotting pathway utilizing factor IX. Such diseases or disease states include stroke, myocardial infarction, aneurysm surgery, and deep vein thrombosis assocd. with surgical procedures, long periods of confinement, and acquired or inherited pro-coagulant states. The pharmaceutical compn. comprising the compd. I is claimed.

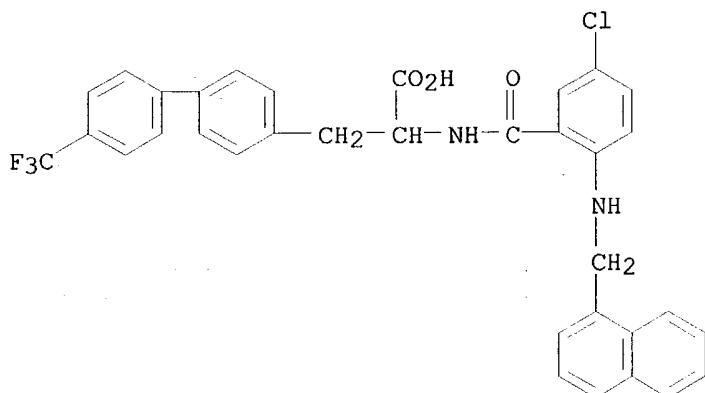
IT 660828-55-7P 660828-57-9P 660828-58-0P
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of substituted (2S)-(arylamino)-3-(biphenyl-4-yl)propionic acids as antagonists of factor IX for inhibiting the intrinsic pathway of blood coagulation)

RN 660828-55-7 CAPLUS

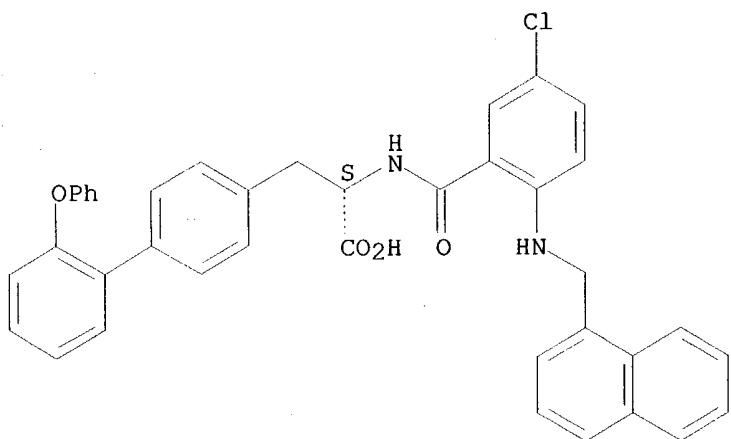
CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-(trifluoromethyl)- (9CI) (CA INDEX NAME)



RN 660828-57-9 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-phenoxy-, (.alpha.S)- (9CI) (CA INDEX NAME)

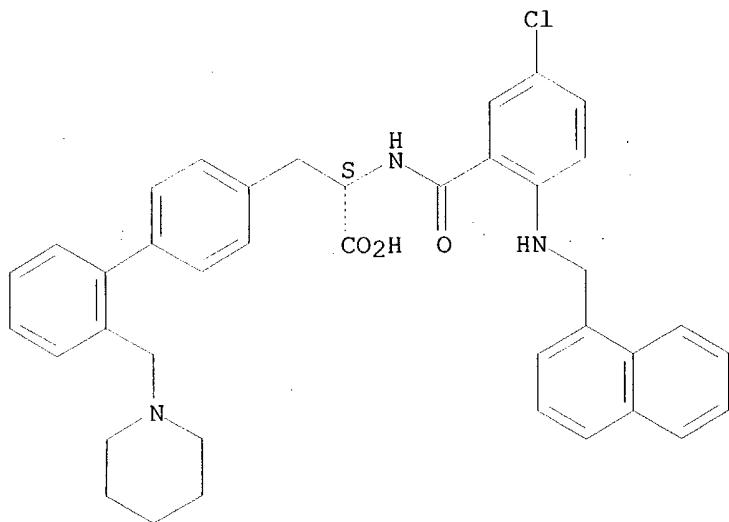
Absolute stereochemistry.



RN 660828-58-0 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-(1-piperidinylmethyl)-, (.alpha.S)- (9CI) (CA INDEX NAME)

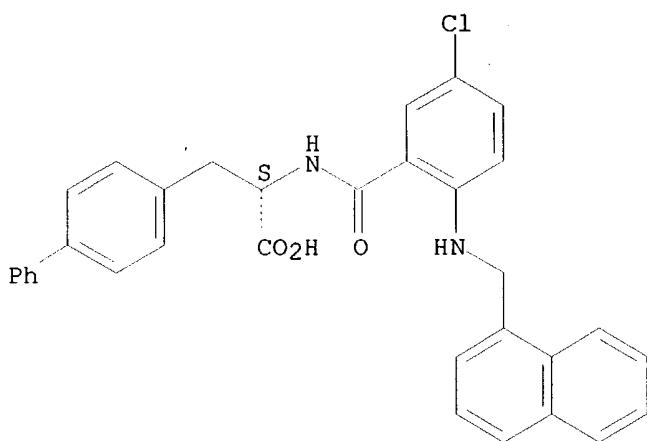
Absolute stereochemistry.



RN 660828-60-4 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-, (.alpha.S)- (9CI) (CA INDEX NAME)

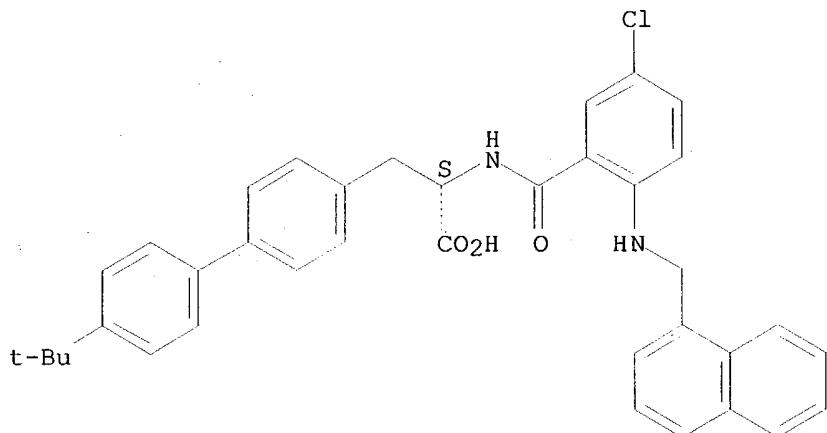
Absolute stereochemistry.



RN 660828-61-5 CAPLUS

RR: 1,1'-Biphenyl-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-(1,1-dimethylethyl)-, (.alpha.S)- (9CI) (CA INDEX NAME)

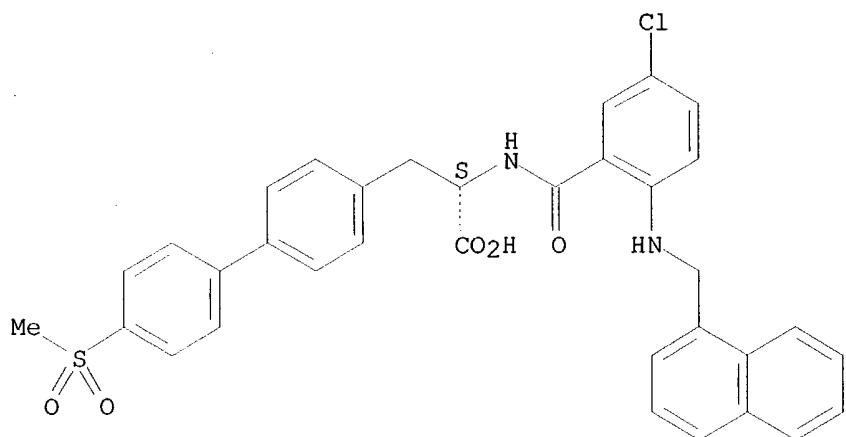
Absolute stereochemistry.



RN 660828-62-6 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-(methylsulfonyl)-, (.alpha.S)-(9CI) (CA INDEX NAME)

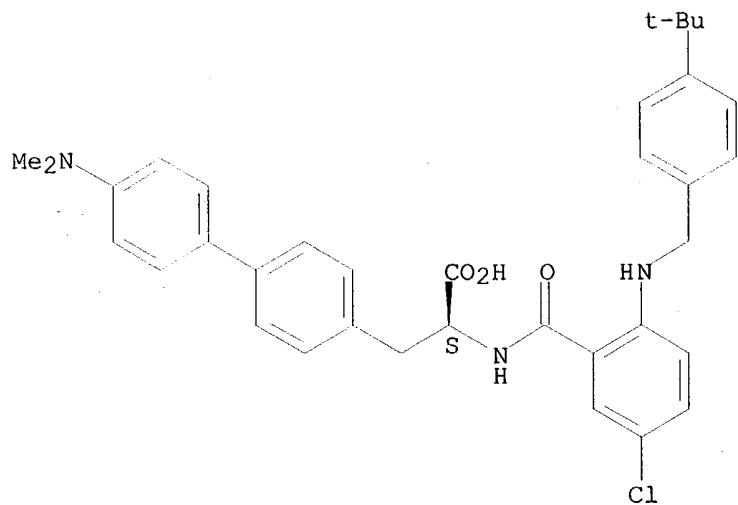
Absolute stereochemistry.



RN 660828-65-9 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-4'-(dimethylamino)-, (.alpha.S)- (9CI) (CA INDEX NAME)

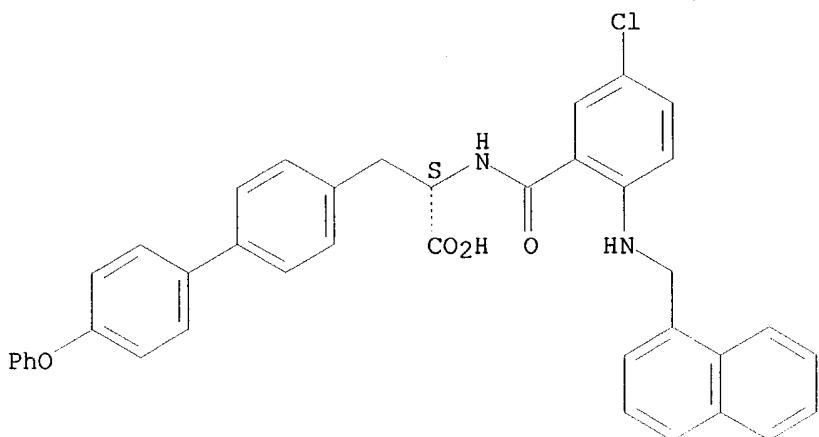
Absolute stereochemistry.



RN 660828-67-1 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-phenoxy-, (.alpha.S)- (9CI) (CA INDEX NAME)

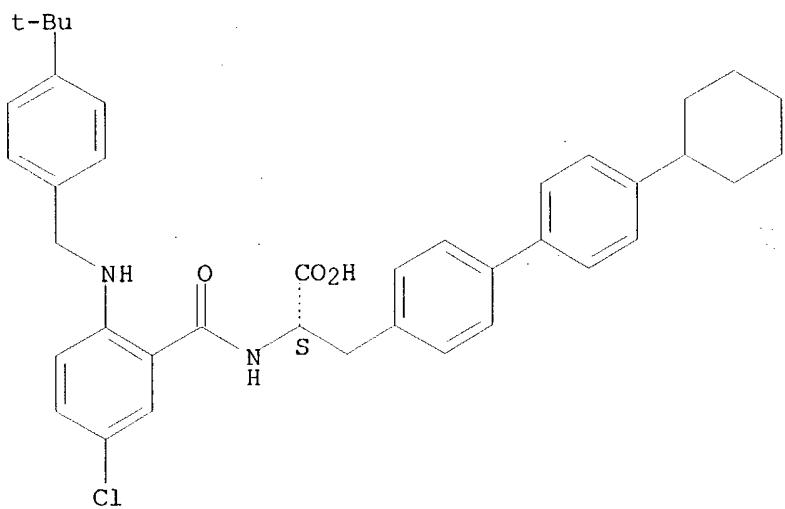
Absolute stereochemistry.



RN 660828-68-2 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-4'-cyclohexyl-, (.alpha.S)- (9CI) (CA INDEX NAME)

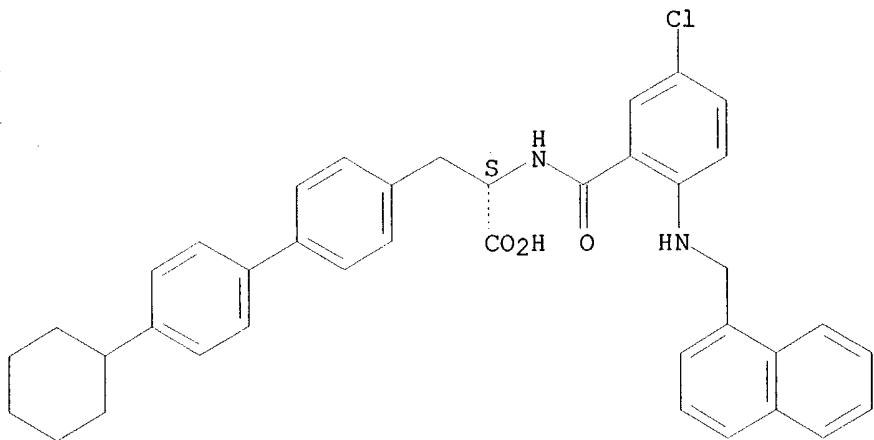
Absolute stereochemistry.



RN 660828-71-7 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-cyclohexyl-, (.alpha.S)- (9CI) (CA INDEX NAME)

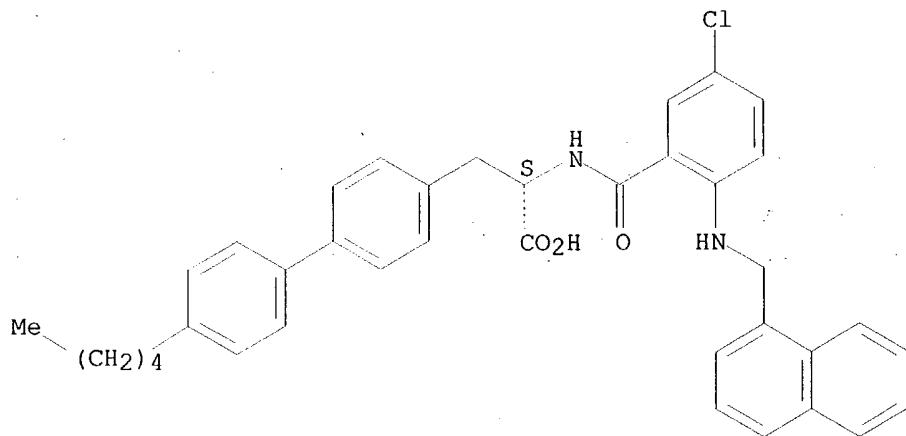
Absolute stereochemistry.



RN 660828-72-8 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, α -[(5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl)amino]-4'-pentyl-, (α .S)- (9CI) (CA INDEX NAME)

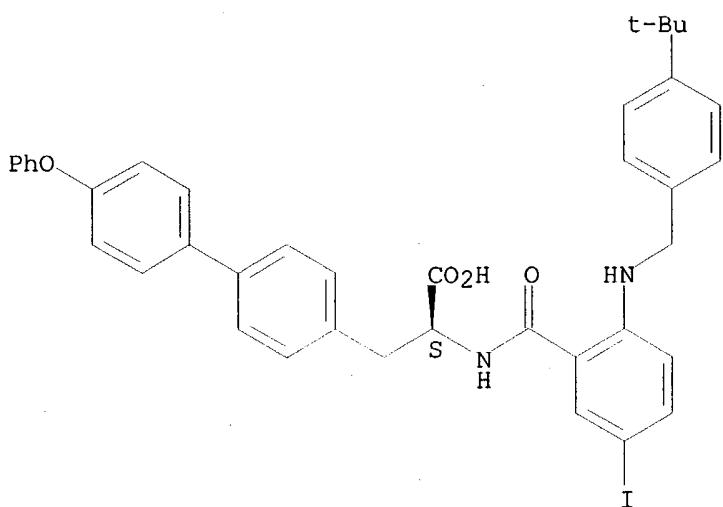
Absolute stereochemistry.



RN 660828-73-9 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, α -[(2-[[4-(1,1-dimethylethyl)phenyl]methyl]amino)-5-iodobenzoyl]amino]-4'-phenoxy-, (α .S)- (9CI) (CA INDEX NAME)

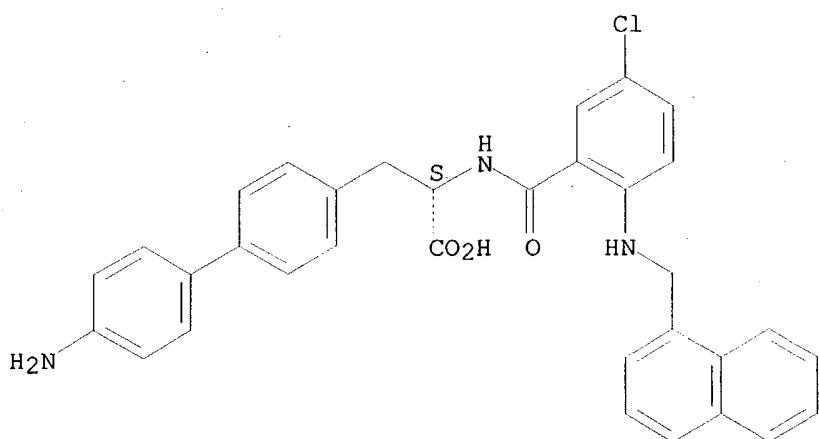
Absolute stereochemistry.



RN 660828-74-0 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, 4'-amino-.alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-, (.alpha.S)- (9CI) (CA INDEX NAME)

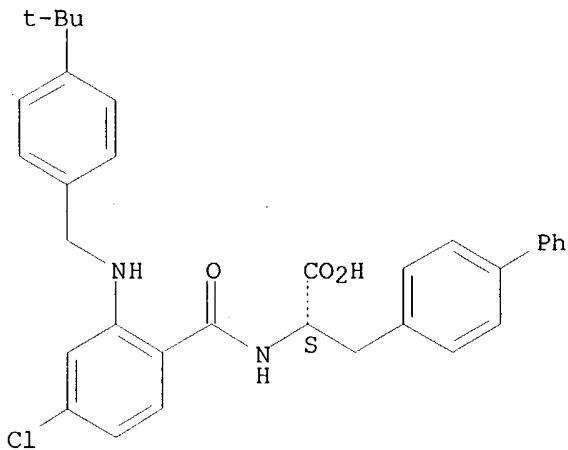
Absolute stereochemistry.



RN 660828-79-5 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[4-chloro-2-[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-, (.alpha.S)- (9CI) (CA INDEX NAME)

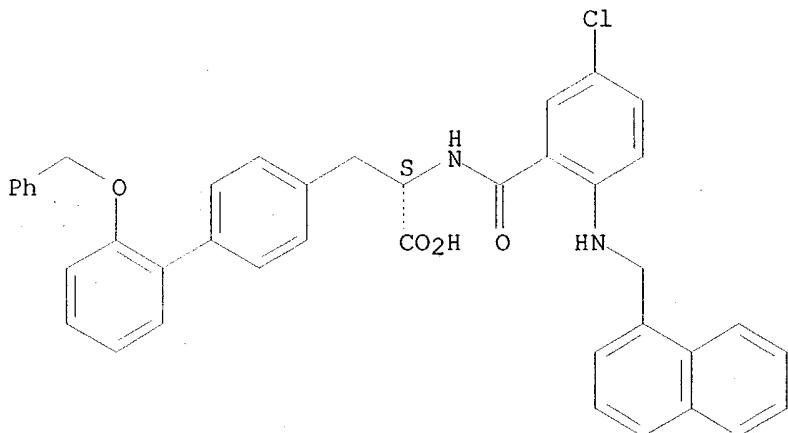
Absolute stereochemistry.



RN 660828-83-1 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-(phenylmethoxy)-, (.alpha.S)-(9CI) (CA INDEX NAME)

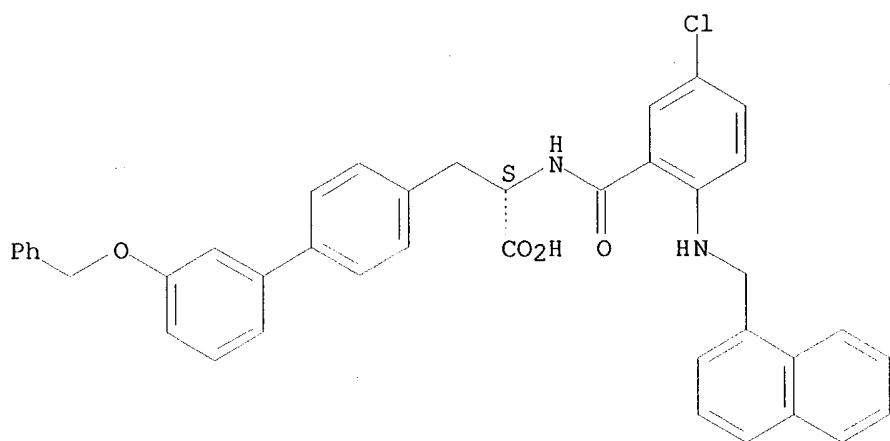
Absolute stereochemistry.



RN 660828-84-2 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-3'-(phenylmethoxy)-, (.alpha.S)-(9CI) (CA INDEX NAME)

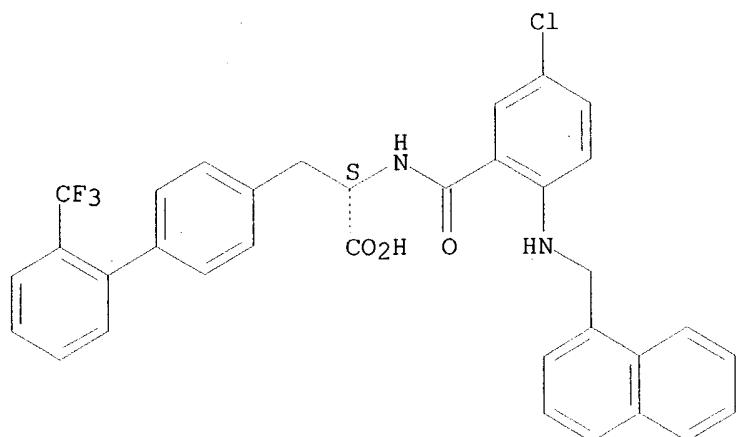
Absolute stereochemistry.



RN 660828-85-3 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-(trifluoromethyl)-, (.alpha.S)- (9CI) (CA INDEX NAME)

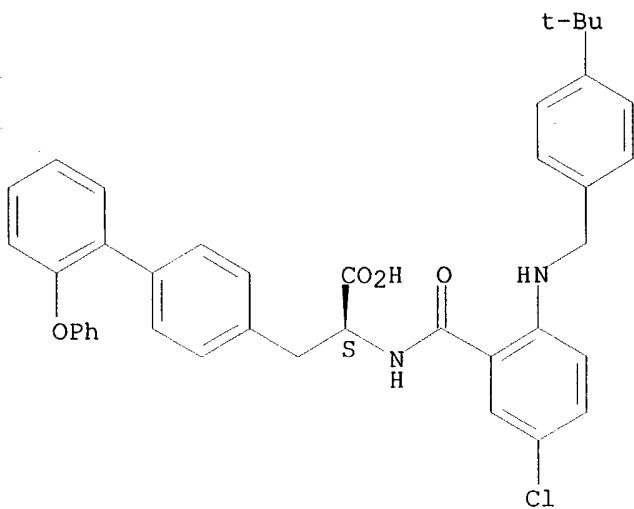
Absolute stereochemistry.



RN 660828-87-5 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-2'-(trifluoromethyl)-, (.alpha.S)- (9CI) (CA INDEX NAME)

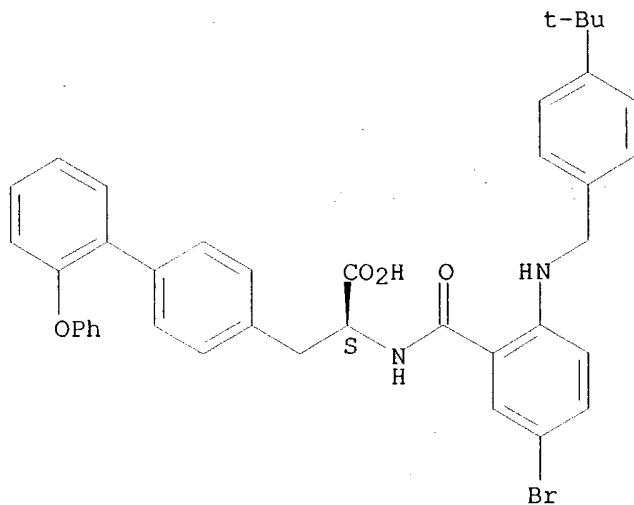
Absolute stereochemistry.



RN 660828-88-6 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-bromo-2-[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-2'-phenoxy-, (.alpha.S)- (9CI) (CA INDEX NAME)

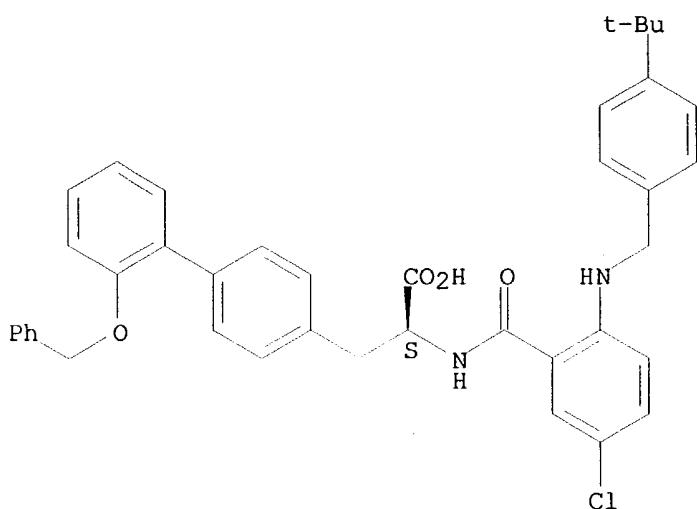
Absolute stereochemistry.



RN 660828-92-2 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[[4-(1,1-dimethylethyl)phenyl]methyl]amino]benzoyl]amino]-2'-(phenylmethoxy)-, (.alpha.S)- (9CI) (CA INDEX NAME)

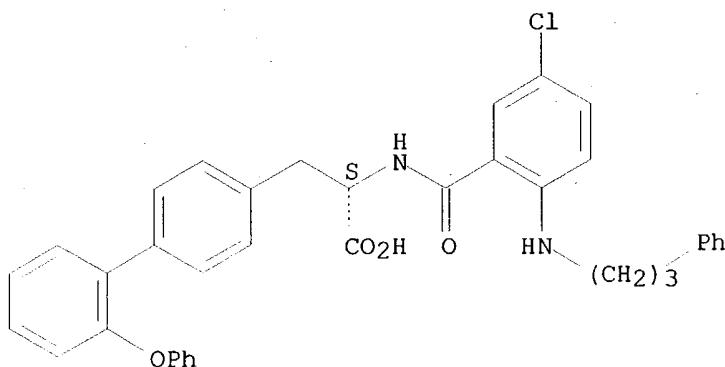
Absolute stereochemistry.



RN 660829-05-0 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(3-phenylpropyl)amino]benzoyl]amino]-2'-phenoxy-, (.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



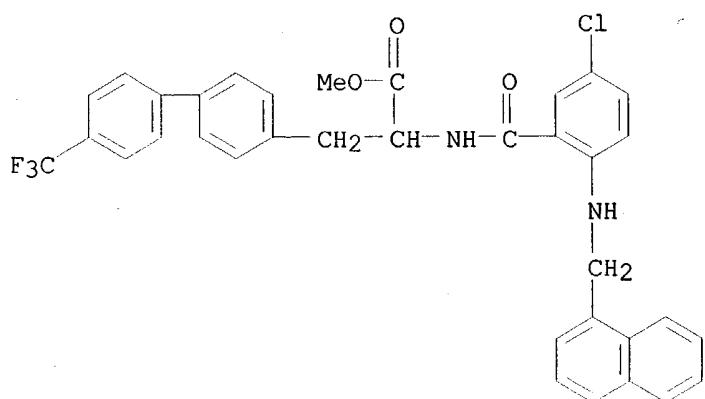
IT 660831-01-6P 660831-06-1P 660831-07-2P

660831-08-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of substituted (2S)- (arylamino)-3- (biphenyl-4-yl)propionic acids as antagonists of factor IX for inhibiting the intrinsic pathway of blood coagulation)

RN 660831-01-6 CAPLUS

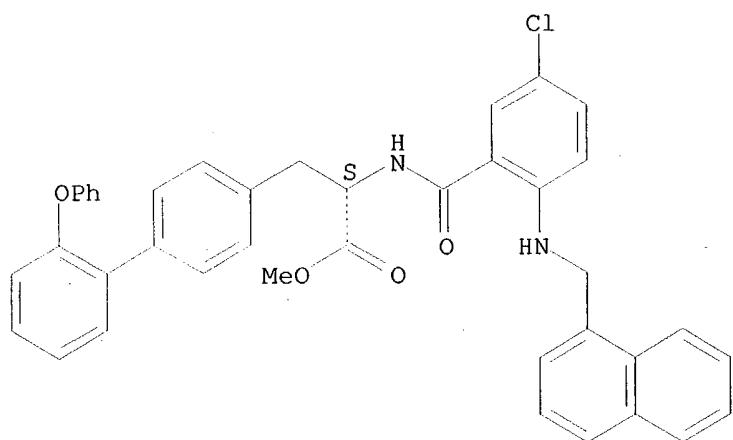
CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-4'-(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 660831-06-1 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-phenoxy-, methyl ester, (.alpha.S)- (9CI) (CA INDEX NAME)

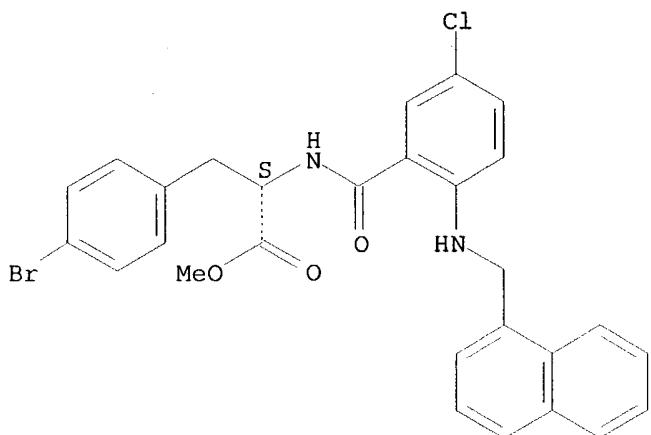
Absolute stereochemistry.



RN 660831-07-2 CAPLUS

CN L-Phenylalanine, 4-bromo-N-[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]-, methyl ester (9CI) (CA INDEX NAME)

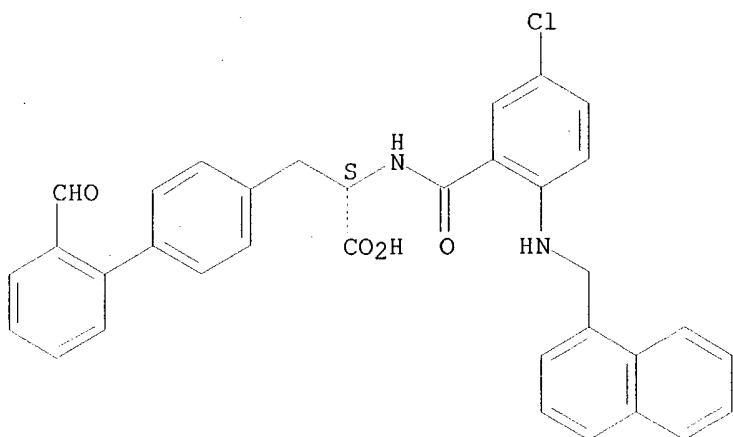
Absolute stereochemistry.



RN 660831-08-3 CAPLUS

CN [1,1'-Biphenyl]-4-propanoic acid, .alpha.-[[5-chloro-2-[(1-naphthalenylmethyl)amino]benzoyl]amino]-2'-formyl-, (.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



✓ L01 ANSWER 4 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:120827 CAPLUS

DOCUMENT NUMBER: 140:181330

TITLE: Preparation of anthranylamidopyridines as inhibitors of vascular endothelial growth factor receptor-2 and -3 (VEGFR-2 and -3).

INVENTOR(S): Huth, Andreas; Krueger, Martin; Zorn, Ludwig; Ince, Stuart; Thierauch, Karl-Heinz; Menrad, Andreas;

PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 88 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

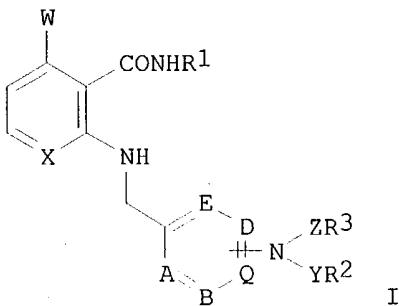
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2004013102	A1	20040212	WO 2003-EP7964	20030722
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

DE 10235690 A1 20040219 DE 2002-10235690 20020731

PRIORITY APPLN. INFO.: DE 2002-10235690 A 20020731
DE 2003-10328036 A 20030619

ED Entered STN: 13 Feb 2004
GI



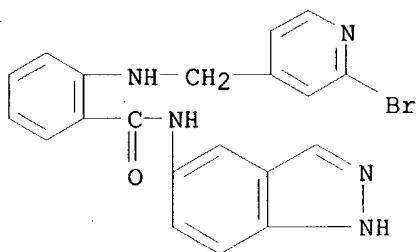
AB Title compds. [I; X = CH, N; W = H, F; A, B, D, E, Q = N, C; .ltoreq.2 of A, B, D, E, Q = N; R1 = (substituted) aryl, heteroaryl; Y, Z = bond, CO, CS, SO2; R2, R3 = H, CONR9R10, SO2R6, COR11, NR9R10, (substituted) alkyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; R2YNZAR3 = atoms to form a 3-8 membered (substituted) (unsatd.) ring; R6 = H, alkyl, haloalkyl, (substituted) aryl, heteroaryl, NR9R10; R9, R10 = H, alkyl, alkenyl, aryl, cycloalkyl, etc.; R11 = alkyl, alkoxy, hydroxyalkyl, hydroxyalkoxy, cycloalkyl, (substituted) Ph, pyridyl, biphenyl, naphthyl], were prep'd. Thus, 2-[(2-bromopyridin-4-ylmethyl)amino]-N-(3-trifluoromethylphenyl)benzamide (prepn. given) pyridine, and N,N-dimethylaminoethylamine were heated in a pressure vessel for 5 h at 200.degree. to give 2-[(2-dimethylaminoethylamino)pyridin-4-ylmethyl]amino]-N-(3-trifluoromethylphenyl)benzamide. I inhibited VEGFR-2 with IC50 = 8-65 nM. I can be used for treatment of tumor or metastasis growth, psoriasis, Kaposi's sarcoma, restenosis, endometriosis, Crohn's disease, Hodgkin's disease, leukemia, arthritis, hemangioma, angiofibroma, eye disease, renal diseases, transplant rejection, fibrotic diseases, mesangial cell proliferative diseases, atherosclerosis, injuries to nervous tissue and for inhibition of the reocclusion of vessels after balloon catheter treatment, in vessel prosthetics, or after the application of mech. devices to hold open vessels, as immunosuppressants, for scar-free wound healing, age spots and contact dermatitis.

IT 657401-05-3 657401-06-4

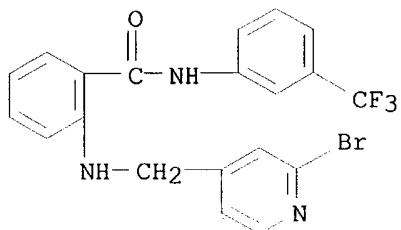
RL: RCT (Reactant); RACT (Reactant or reagent)
(prepn. of anthranylamidopyridines as inhibitors of vascular endothelial growth factor receptor)

RN 657401-05-3 CAPLUS

CN Benzamide, 2-[(2-bromo-4-pyridinyl)methyl]amino]-N-1H-indazol-5-yl- (9CI)
(CA INDEX NAME)

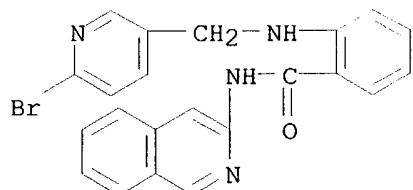


RN 657401-06-4 CAPLUS
CN Benzamide, 2-[(2-bromo-4-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

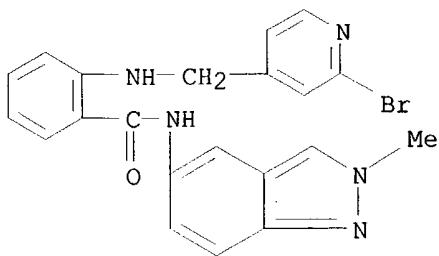


IT 474799-36-5P 657401-01-9P 657401-04-2P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of anthranyl amidopyridines as inhibitors of vascular endothelial growth factor receptor)

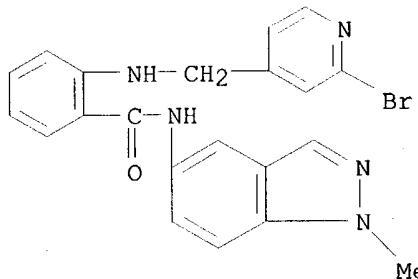
RN 474799-36-5 CAPLUS
CN Benzamide, 2-[(6-bromo-3-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI)
(CA INDEX NAME)



RN 657401-01-9 CAPLUS
CN Benzamide, 2-[(2-bromo-4-pyridinyl)methyl]amino]-N-(2-methyl-2H-indazol-5-yl)- (9CI) (CA INDEX NAME)



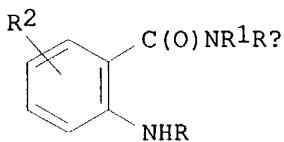
RN 657401-04-2 CAPLUS
 CN Benzamide, 2-[(2-bromo-4-pyridinyl)methyl]amino]-N-(1-methyl-1H-indazol-5-yl)- (9CI) (CA INDEX NAME)



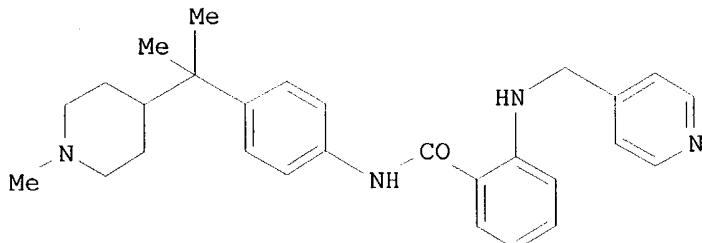
ANSWER 5 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2004:41461 CAPLUS
 DOCUMENT NUMBER: 140:93789
 TITLE: Preparation of substituted anthranilic amide derivatives as VEGF modulators and methods of use against cancer and other disorders
 INVENTOR(S): Huang, Qi; Chen, Guoqing; Li, Aiwen; Riahi, Babak; Tasker, Andrew; Yang, Kevin; Yuan, Chester Chenguang
 PATENT ASSIGNEE(S): Amgen Inc., USA
 SOURCE: PCT Int. Appl., 204 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004005279	A2	20040115	WO 2003-US21601	20030709
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
PRIORITY APPLN. INFO.:			US 2002-395144P	P 20020709
			US 2003-615809	A 20030708
OTHER SOURCE(S):		MARPAT 140:93789		

ED Entered STN: 18 Jan 2004
 GI



I



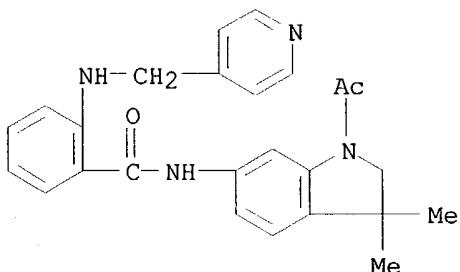
II

AB Selected substituted anthranilic amide derivs. (shown as I; variables defined below; e.g. II) are effective for prophylaxis and treatment of diseases, such as angiogenesis mediated diseases. The invention encompasses novel compds., analogs, prodrugs and pharmaceutically acceptable salts thereof, pharmaceutical compns. and methods for prophylaxis and treatment of diseases and other maladies or conditions involving cancer and the like. Although the methods of prepn. are not claimed, .apprx.139 example prepn. of I and .apprx.80 of intermediates are included. For example, II was prepnd. in 3 steps starting from 2-nitrobenzoic acid and [4-[1-Methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]amine and involving intermediates 2-nitro-N-[4-[1-methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]benzamide and 2-amino-N-[4-[1-methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]benzamide. Compds. I showed inhibition of KDR at doses <50 .mu.M. Some of the exemplified I inhibit VEGF-stimulated HUVEC proliferation <1 .mu.M. Compds. I are active at doses <150 mpk in a tumor model. For I: R = (un)substituted 9- or 10-membered fused heterocyclyl, -(CH₂)₁₋₂-R₃; R₁ = (un)substituted 5-6 membered satd. or partially satd. heterocyclyl, 9-10 membered bicyclic and 13-14 membered tricyclic satd. or partially satd. heterocyclyl, and phenyl; R₂ is .gtoreq.1 substituents = H, halo, hydroxy, amino, C1-6-alkyl, C1-6-haloalkyl, C1-6-alkoxy, C1-2-alkylamino, aminosulfonyl, C3-6-cycloalkyl, cyano, C1-2-hydroxyalkyl, nitro, C2-3-alkenyl, C2-3-alkynyl, C1-6-haloalkoxy, C1-6-carboxyalkyl, 4-6-membered heterocyclyl-C1-6-alkylamino, (un)substituted Ph and (un)substituted 4-6 membered heterocyclyl; Ra = H, C1-2-alkyl; addnl. details are given in the claims.

IT **645418-47-9P**, N-(1-Acetyl-3,3-dimethyl-2,3-dihydro-1H-indol-6-yl)-2-[(pyridin-4-yl)methyl]amino]benzamide **645418-59-3P**, N-(1-Acetyl-3,3-dimethyl-2,3-dihydro-1H-indol-6-yl)-2-[(quinolin-4-yl)methyl]amino]benzamide
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
 (drug candidate; prepn. of substituted anthranilic amide derivs. as VEGF modulators and methods of use against cancer and other disorders)

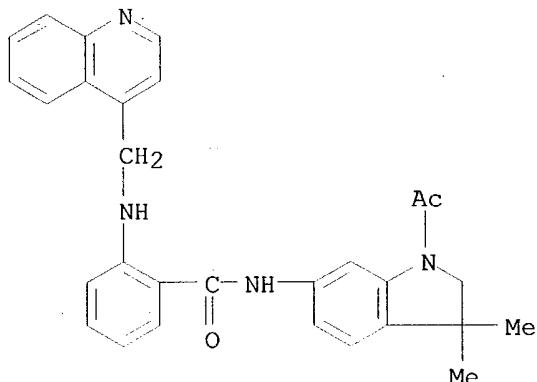
RN 645418-47-9 CAPLUS

CN Benzamide, N-(1-acetyl-2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 645418-59-3 CAPLUS

CN Benzamide, N-(1-acetyl-2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)



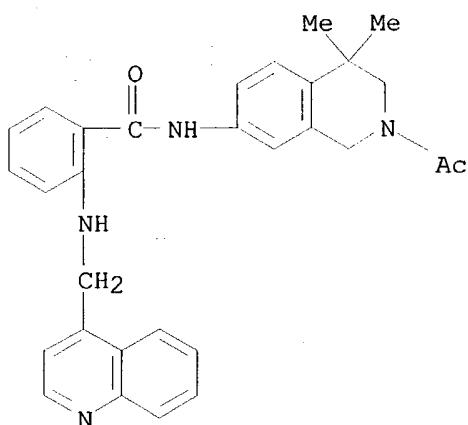
IT 453564-10-8P, N-(2-Acetyl-4,4-dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-[(quinolin-4-yl)methyl]amino]benzamide
 645418-43-5P, N-[4-[1-Methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]-2-[(pyridin-4-yl)methyl]amino]benzamide
 645418-48-0P, N-(1-Acetyl-3,3-dimethyl-2,3-dihydro-1H-indol-6-yl)-2-(4-fluorobenzylamino)benzamide 645418-49-1P,
 N-[4-[1-Methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]-2-[(quinolin-4-yl)methyl]amino]benzamide 645418-50-4P, 2-(4-Fluorobenzylamino)-
 N-[4-[1-methyl-1-(1-methylpiperidin-4-yl)ethyl]phenyl]benzamide
 645418-51-5P, N-(3,3-Dimethyl-2,3-dihydro-1H-indol-6-yl)-2-[(pyridin-4-yl)methyl]amino]benzamide 645418-52-6P,
 N-(1-Ethyl-3,3-dimethyl-2,3-dihydro-1H-indol-6-yl)-2-[(pyridin-4-yl)methyl]amino]benzamide 645418-56-0P, N-[3,3-Dimethyl-1-[(4-methylpiperazin-1-yl)carbonyl]-2,3-dihydro-1H-indol-6-yl]-2-(4-fluorobenzylamino)benzamide 645418-62-8P, N-(3,3-Dimethyl-2,3-dihydro-1H-indol-6-yl)-2-[(quinolin-4-yl)methyl]amino]benzamide
 645418-64-0P, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-(4-fluorobenzylamino)benzamide 645418-67-3P,
 N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-fluoro-6-(4-fluorobenzylamino)benzamide 645418-68-4P, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-3-fluoro-6-(4-fluorobenzylamino)benzamide 645418-69-5P, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-4-fluoro-6-(4-fluorobenzylamino)benzamide 645418-70-8P, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-3,4-difluoro-6-(4-fluorobenzylamino)benzamide 645418-71-9P, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-[(2-methoxypyridin-4-yl)methyl]amino]benzamide 645418-74-2P, N-(4,4-Dimethyl-1,2,3,4-

tetrahydroisoquinolin-7-yl)-2-fluoro-6-[(2-methoxypyridin-4-yl)methyl]amino]benzamide **645418-75-3P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-3-fluoro-6-[(2-methoxypyridin-4-yl)methyl]amino]benzamide **645418-76-4P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-4-fluoro-6-[(2-methoxypyridin-4-yl)methyl]amino]benzamide **645418-78-6P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-[(1H-pyrrolo[2,3-b]pyridin-3-yl)methyl]amino]benzamide **645418-81-1P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroquinolin-7-yl)-2-[(1H-pyrrolo[2,3-b]pyridin-3-yl)methyl]amino]benzamide **645418-82-2P**, N-(4,4-Dimethyl-2-oxo-1,2,3,4-tetrahydroquinolin-7-yl)-2-[(1H-pyrrolo[2,3-b]pyridin-3-yl)methyl]amino]benzamide **645418-83-3P**, 1,1-Dimethylethyl 7-[[2-[(7-fluoro-1H-indol-3-yl)methyl]amino]phenyl]carbonyl]amino]-4,4-dimethyl-3,4-dihydro-2(1H)-isoquinolinecarboxylate **645418-97-9P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-[(pyridazin-4-yl)methyl]amino]benzamide **645418-98-0P**, 4,4-Dimethyl-7-[[2-[(quinoxalin-5-yl)methyl]amino]benzoyl]amino]-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester **645418-99-1P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-2-[(quinoxalin-5-yl)methyl]amino]benzamide **645419-14-3P**, N-(4,4-Dimethyl-1,2,3,4-tetrahydroisoquinolin-7-yl)-3-fluoro-2-(4-fluorobenzylamino)benzamide
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; prepn. of substituted anthranilic amide derivs. as VEGF modulators and methods of use against cancer and other disorders)

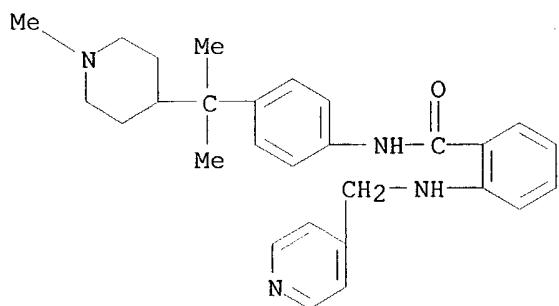
RN 453564-10-8 CAPLUS

CN Benzamide, N-(2-acetyl-1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinoliny1)-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)



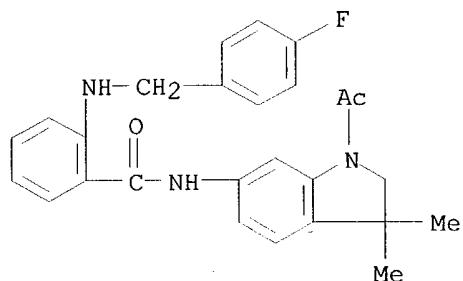
RN 645418-43-5 CAPLUS

CN Benzamide, N-[4-[1-methyl-1-(1-methyl-4-piperidinyl)ethyl]phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



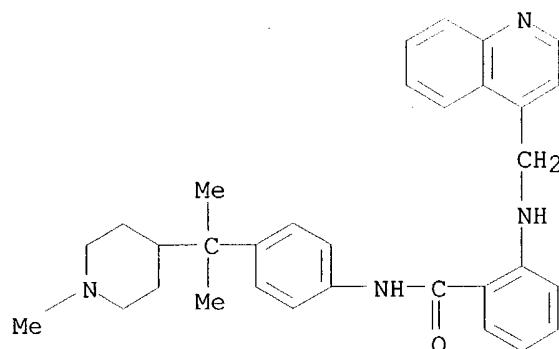
RN 645418-48-0 CAPLUS

CN Benzamide, N-(1-acetyl-2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-fluorophenyl)methyl]amino]- (9CI) (CA INDEX NAME)



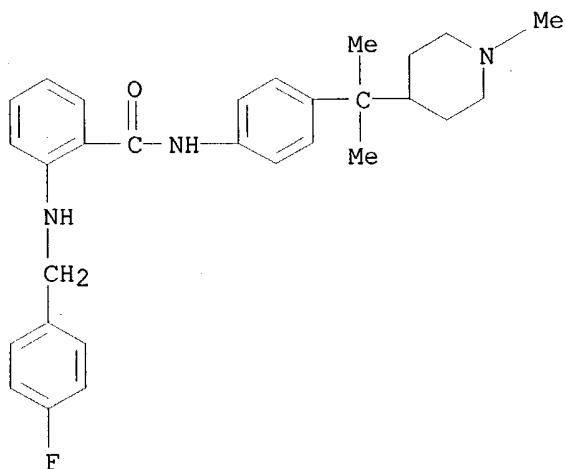
RN 645418-49-1 CAPLUS

CN Benzamide, N-[4-[1-methyl-1-(1-methyl-4-piperidinyl)ethyl]phenyl]-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)

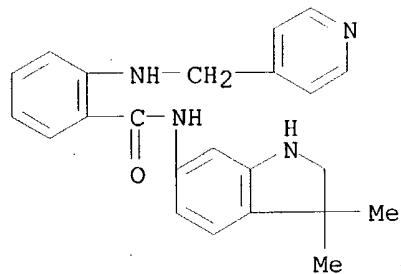


RN 645418-50-4 CAPLUS

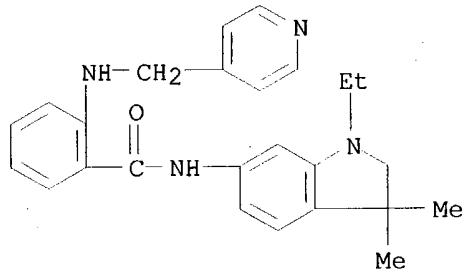
CN Benzamide, 2-[(4-fluorophenyl)methyl]amino]-N-[4-[1-methyl-1-(1-methyl-4-piperidinyl)ethyl]phenyl]- (9CI) (CA INDEX NAME)



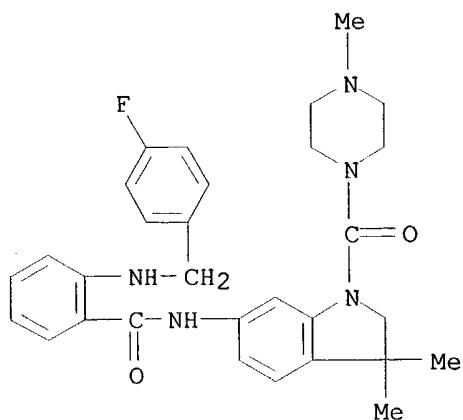
RN 645418-51-5 CAPLUS
CN Benzamide, N-(2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



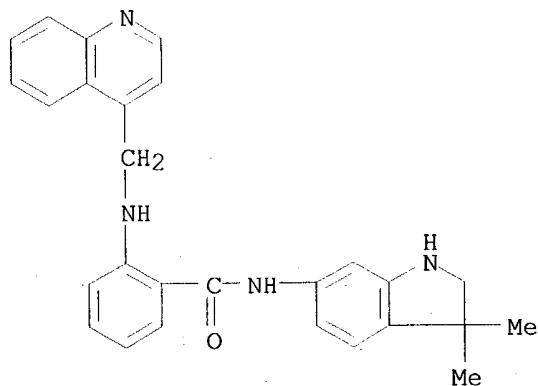
RN 645418-52-6 CAPLUS
CN Benzamide, N-(1-ethyl-2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



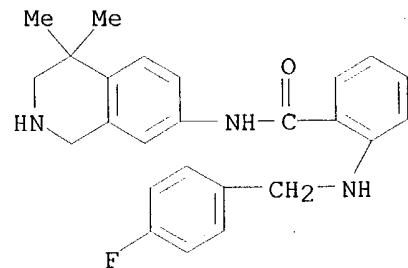
RN 645418-56-0 CAPLUS
CN Benzamide, N-[2,3-dihydro-3,3-dimethyl-1-[(4-methyl-1-piperazinyl)carbonyl]-1H-indol-6-yl]-2-[(4-fluorophenyl)methyl]amino]- (9CI) (CA INDEX NAME)



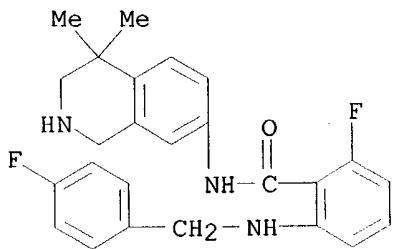
RN 645418-62-8 CAPLUS
 CN Benzamide, N-(2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)



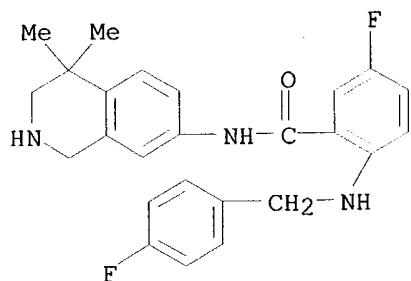
RN 645418-64-0 CAPLUS
 CN Benzamide, 2-[(4-fluorophenyl)methylamino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



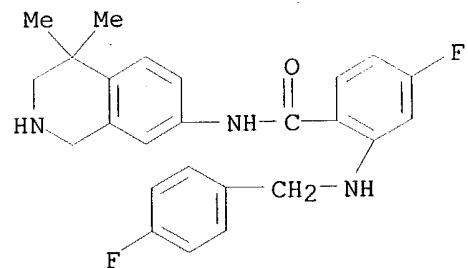
RN 645418-67-3 CAPLUS
 CN Benzamide, 2-fluoro-6-[(4-fluorophenyl)methylamino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



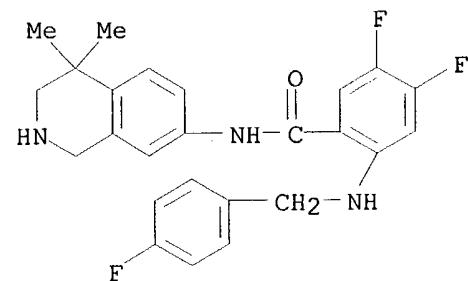
RN 645418-68-4 CAPLUS
CN Benzamide, 5-fluoro-2-[(4-fluorophenyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



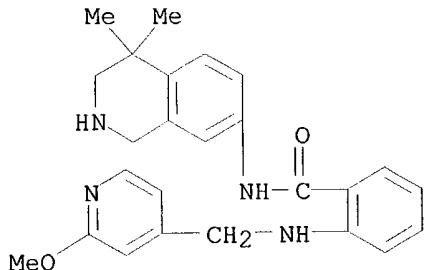
RN 645418-69-5 CAPLUS
CN Benzamide, 4-fluoro-2-[(4-fluorophenyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



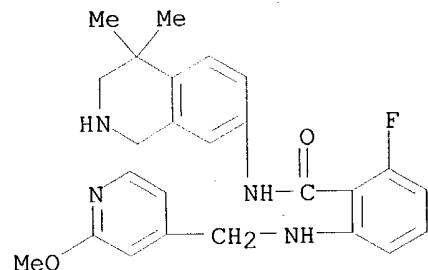
RN 645418-70-8 CAPLUS
CN Benzamide, 4,5-difluoro-2-[(4-fluorophenyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



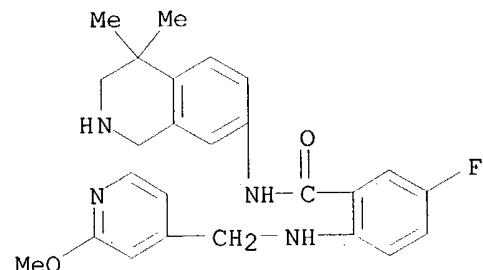
RN 645418-71-9 CAPLUS
CN Benzamide, 2-[(2-methoxy-4-pyridinyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



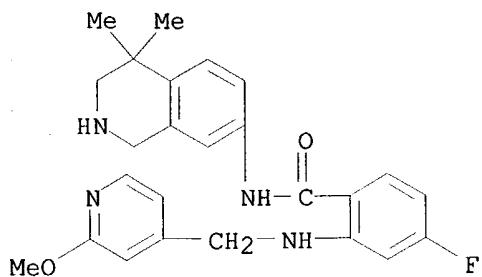
RN 645418-74-2 CAPLUS
CN Benzamide, 2-fluoro-6-[(2-methoxy-4-pyridinyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



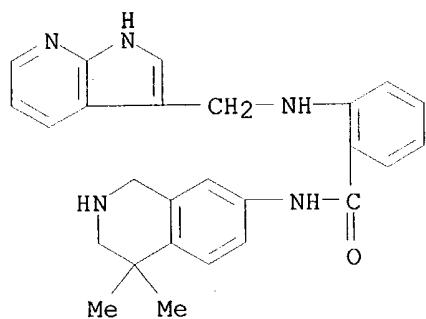
RN 645418-75-3 CAPLUS
CN Benzamide, 5-fluoro-2-[(2-methoxy-4-pyridinyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



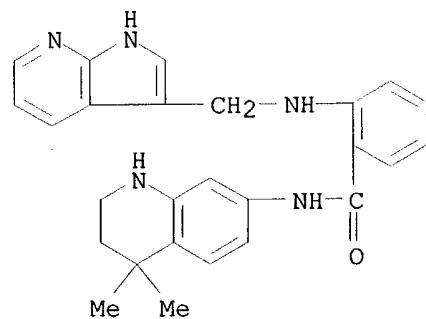
RN 645418-76-4 CAPLUS
CN Benzamide, 4-fluoro-2-[(2-methoxy-4-pyridinyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



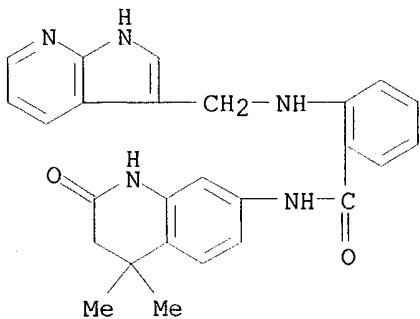
RN 645418-78-6 CAPLUS
 CN Benzamide, 2-[(1H-pyrrolo[2,3-b]pyridin-3-ylmethyl)amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



RN 645418-81-1 CAPLUS
 CN Benzamide, 2-[(1H-pyrrolo[2,3-b]pyridin-3-ylmethyl)amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-quinolinyl)- (9CI) (CA INDEX NAME)

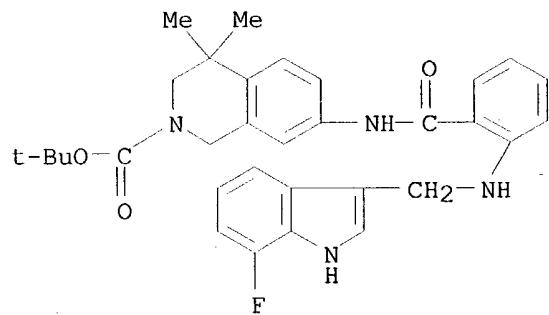


RN 645418-82-2 CAPLUS
 CN Benzamide, 2-[(1H-pyrrolo[2,3-b]pyridin-3-ylmethyl)amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-2-oxo-7-quinolinyl)- (9CI) (CA INDEX NAME)



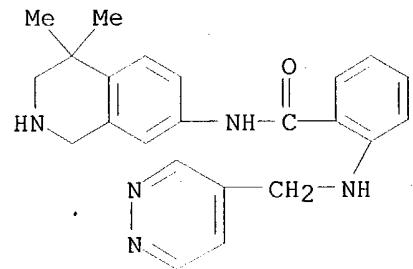
RN 645418-83-3 CAPLUS

CN 2(1H)-Isoquinolinecarboxylic acid, 7-[[2-[(7-fluoro-1H-indol-3-yl)methyl]amino]benzoyl]amino]-3,4-dihydro-4,4-dimethyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



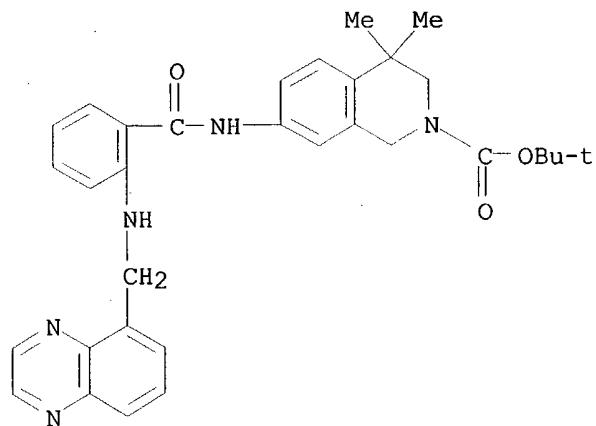
RN 645418-97-9 CAPLUS

CN Benzamide, 2-[(4-pyridazinylmethyl)amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)- (9CI) (CA INDEX NAME)



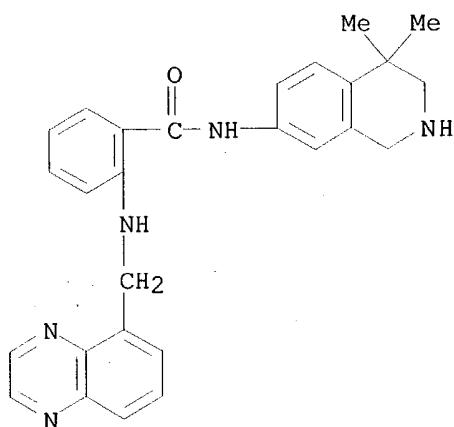
RN 645418-98-0 CAPLUS

CN 2(1H)-Isoquinolinecarboxylic acid, 3,4-dihydro-4,4-dimethyl-7-[[2-[(5-quinoxalinylmethyl)amino]benzoyl]amino]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



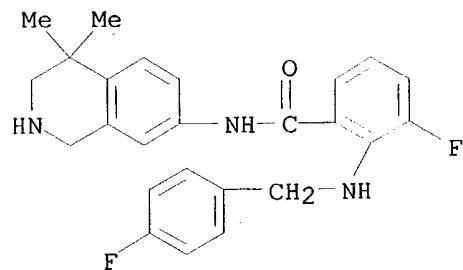
RN 645418-99-1 CAPLUS

CN Benzamide, 2-[(5-quinoxalinylmethyl)amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isooquinolinyl)- (9CI) (CA INDEX NAME)



RN 645419-14-3 CAPLUS

CN Benzamide, 3-fluoro-2-[(4-fluorophenyl)methyl]amino]-N-(1,2,3,4-tetrahydro-4,4-dimethyl-7-isooquinolinyl)- (9CI) (CA INDEX NAME)



IT 645418-65-1P, 7-[(2-(4-Fluorobenzylamino)benzoyl)amino]-4,4-dimethyl-3,4-dihydro-1H-isooquinoline-2-carboxylic acid tert-butyl ester

645418-73-1P, 7-[(2-[(2-Methoxypyridin-4-yl)methyl]amino)benzoyl]amino]-4,4-dimethyl-3,4-dihydro-1H-isooquinoline-2-carboxylic acid tert-butyl ester 645418-80-0P,
4,4-Dimethyl-7-[(2-[(1H-pyrrolo[2,3-b]pyridin-3-

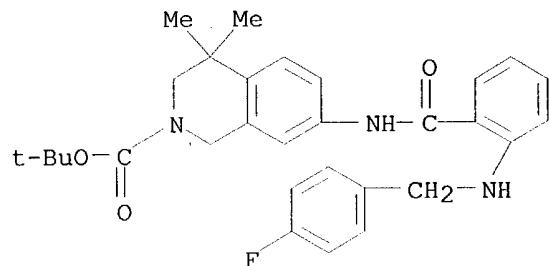
yl)methyl]amino]benzoyl]amino]-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of substituted anthranilic amide derivs. as VEGF modulators and methods of use against cancer and other disorders)

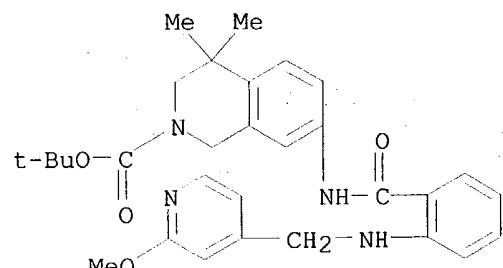
RN 645418-65-1 CAPLUS

CN 2(1H)-Isoquinolinecarboxylic acid, 7-[[2-[[[4-fluorophenyl)methyl]amino]benzoyl]amino]-3,4-dihydro-4,4-dimethyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



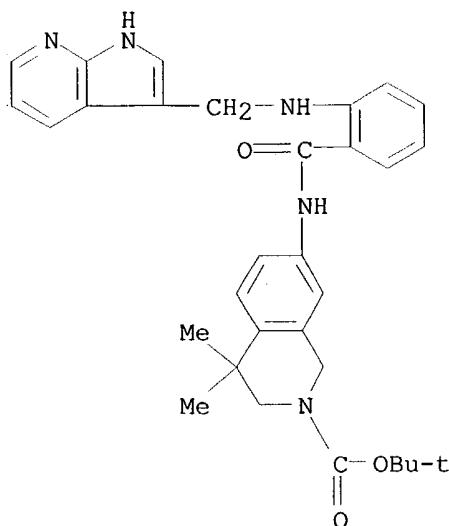
RN 645418-73-1 CAPLUS

CN 2(1H)-Isoquinolinecarboxylic acid, 3,4-dihydro-7-[[2-[[[2-methoxy-4-pyridinyl)methyl]amino]benzoyl]amino]-4,4-dimethyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 645418-80-0 CAPLUS

CN 2(1H)-Isoquinolinecarboxylic acid, 3,4-dihydro-4,4-dimethyl-7-[[2-[(1H-pyrrolo[2,3-b]pyridin-3-ylmethyl)amino]benzoyl]amino]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



ANSWER 6 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:36626 CAPLUS

DOCUMENT NUMBER: 140:93929

TITLE: Preparation of N-(pyridinylmethyl)anthranilamides as VEGFR-2 and VEGFR-3 inhibitors for treating diseases caused by persistent angiogenesis

INVENTOR(S): Huth, Andreas; Krueger, Martin; Zorn, Ludwig; Ince, Stuart; Thierauch, Karl-Heinz; Menrad, Andreas; Haberey, Martin; Hess-Stumpp, Holger

PATENT ASSIGNEE(S): Schering AG, Germany

SOURCE: Ger. Offen., 18 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

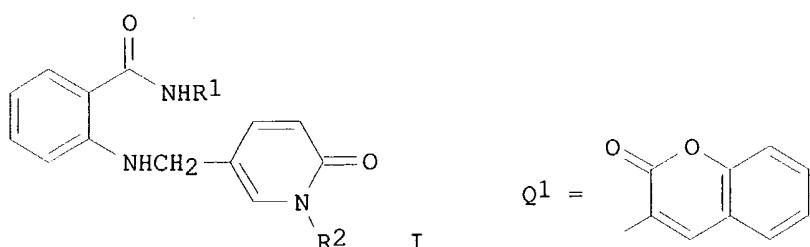
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10228090	A1	20040115	DE 2002-10228090	20020619
US 2004039019	A1	20040226	US 2003-464853	20030619
PRIORITY APPLN. INFO.:			DE 2002-10228090 A	20020619
			US 2002-404773P	P 20020821

OTHER SOURCE(S): MARPAT 140:93929

ED Entered STN: 16 Jan 2004

GI



AB Title compds. [I; R1 = (substituted) indazolyl, indolinyl, quinolinyl, Q1; R2 = H, C1-3 alkyl], were prep'd. Thus, 2-amino-N-(2-oxo-2,3-dihydro-1N-indol-6-yl)benzamide and pyridin-2-one-5-carboxaldehyde in MeOH was treated with ice AcOH followed by stirring over night at room temp. to give 82% N-(2-oxo-2,3-dihydro-1H-indol-6-yl)-2-[(6-oxo-1,6-dihydropyridin-3-yl)methylamino]benzamide. The latter inhibited VEGFR-2 (KDR) with IC50 = 0,05 .mu.M.

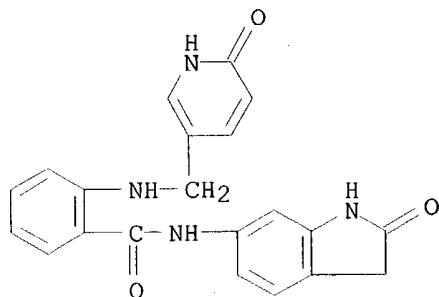
IT 643081-97-4P 643081-98-5P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of N-(pyridinylmethyl)anthranilamides as VEGFR-2 and VEGFR-3 inhibitors for treating diseases caused by persistent angiogenesis)

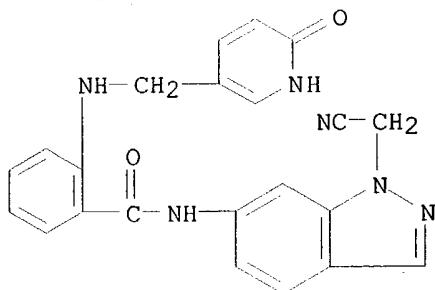
RN 643081-97-4 CAPLUS

CN Benzamide, N-(2,3-dihydro-2-oxo-1H-indol-6-yl)-2-[(1,6-dihydro-6-oxo-3-pyridinyl)methyl]amino]- (9CI) (CA INDEX NAME)



RN 643081-98-5 CAPLUS

CN Benzamide, N-[1-(cyanomethyl)-1H-indazol-6-yl]-2-[(1,6-dihydro-6-oxo-3-pyridinyl)methyl]amino]- (9CI) (CA INDEX NAME)



L31 ANSWER 7 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2003:551370 CAPLUS

DOCUMENT NUMBER: 139:111679

TITLE: Combination of microsomal triglyceride transfer protein (MTP) inhibitors or apoB secretion inhibitors with fibrates for use as drugs

INVENTOR(S): Thomas, Leo; Mark, Michael

PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G., Germany

SOURCE: PCT Int. Appl., 220 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003057205	A2	20030717	WO 2003-EP57	20030107
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
DE 10200633	A1	20030724	DE 2002-10200633	20020110
US 2003162788	A1	20030828	US 2003-339088	20030109
PRIORITY APPLN. INFO.:			DE 2002-10200633 A	20020110
			DE 2002-10256184 A	20021202
			US 2002-353397P	20020201
			US 2002-435386P	20021220

OTHER SOURCE(S): MARPAT 139:111679

ED Entered STN: 18 Jul 2003

AB The invention discloses the use of fibrates for reducing the hepatic toxicity of MTP inhibitors, as well as pharmaceutical compns. that contain an MTP inhibitor and a fibrate. Compd. prepn. is included.

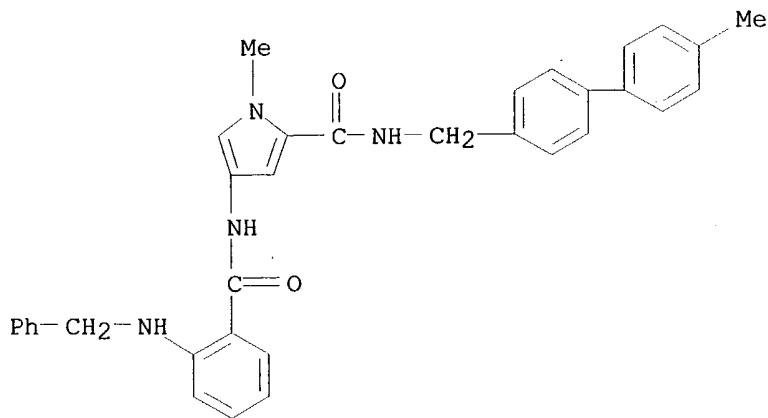
IT 486436-62-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(combination of microsomal triglyceride transfer protein inhibitors or apoB secretion inhibitors with fibrates for use as drugs)

RN 486436-62-8 CAPLUS

CN 1H-Pyrrole-2-carboxamide, 1-methyl-N-[(4'-methyl[1,1'-biphenyl]-4-yl)methyl]-4-[(2-[(phenylmethyl)amino]benzoyl)amino]- (9CI) (CA INDEX NAME)



ANSWER 8 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2003:376825 CAPLUS
 DOCUMENT NUMBER: 138:385308
 TITLE: Preparation of anthranilic acid amides and their use as vascular endothelial growth factor receptor tyrosine kinase inhibitors

INVENTOR(S): Bold, Guido; Furet, Pascal; Manley, Paul William
 PATENT ASSIGNEE(S): Novartis AG, Switz.; Novartis Pharma GmbH
 SOURCE: PCT Int. Appl., 31 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

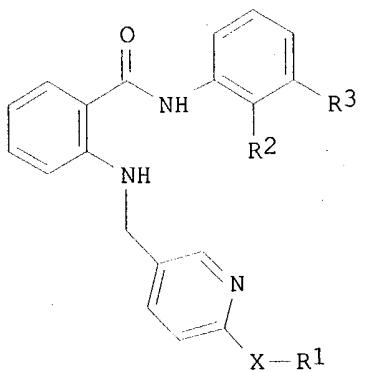
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003040102	A1	20030515	WO 2002-EP12444	20021107
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LT, LU, LV, MA, MD, MK, MN, MX, NO, NZ, OM, PH, PL, PT, RO, RU, SE, SG, SI, SK, TJ, TM, TN, TR, TT, UA, US, UZ, VC, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR				

PRIORITY APPLN. INFO.: GB 2001-26902 A 20011108

OTHER SOURCE(S): MARPAT 138:385308

ED Entered STN: 16 May 2003

GI



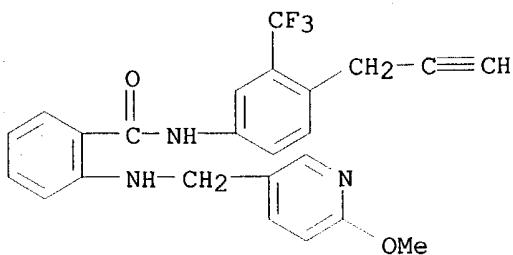
AB Anthranilic acid amide derivs. [I; R1, R2 = H, lower alkyl; R3 = lower perfluoroalkyl; X = O, S; e.g., 2-[(6-Methoxy-3-pyridinyl)methyl]amino-N-[3-(trifluoromethyl)phenyl]benzamide hydrochloride, m.p. 133-135.degree.], which are vascular endothelial growth factor receptor tyrosine kinase inhibitors for the treatment of neoplastic disease, of retinopathy or age-related macular degeneration, are prepd. and a I-contg. formulation presented (e.g., a soft capsule).

IT 524941-34-2

RL: RCT (Reactant); RACT (Reactant or reagent)
 (in the prepn. of anthranilic acid amides)

RN 524941-34-2 CAPLUS

CN Benzamide, 2-[[[(6-methoxy-3-pyridinyl)methyl]amino]-N-[4-(2-propynyl)-3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

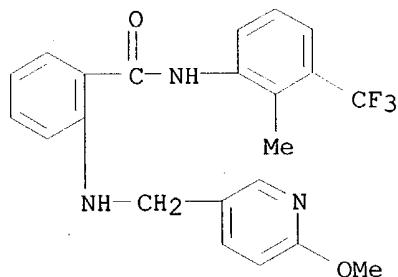


IT 524941-29-5P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(in the prepn. of anthranilic acid amides for use as vascular endothelial growth factor receptor tyrosine kinase inhibitors)

RN 524941-29-5 CAPLUS

CN Benzamide, 2-[(6-methoxy-3-pyridinyl)methylamino]-N-[2-methyl-3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

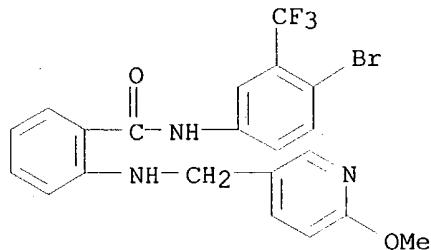


IT 524728-97-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of)

RN 524728-97-0 CAPLUS

CN Benzamide, N-[4-bromo-3-(trifluoromethyl)phenyl]-2-[(6-methoxy-3-pyridinyl)methylamino]- (9CI) (CA INDEX NAME)

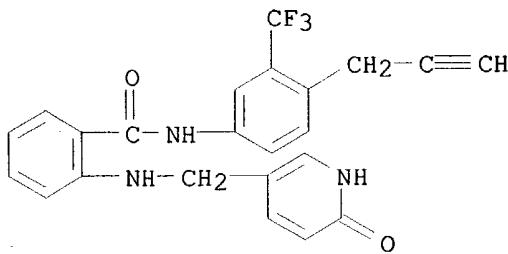


IT 524941-33-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 524941-33-1 CAPLUS

CN Benzamide, 2-[(1,6-dihydro-6-oxo-3-pyridinyl)methylamino]-N-[4-(2-propynyl)-3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



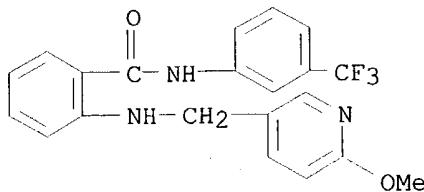
IT 524941-28-4P

RL: RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use);
 BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent);
 USES (Uses)

(prepn. of anthranilic acid amides and their use as vascular
 endothelial growth factor receptor tyrosine kinase inhibitors)

RN 524941-28-4 CAPLUS

CN Benzamide, 2-[(6-methoxy-3-pyridinyl)methyl]amino-N-[3-(trifluoromethyl)phenyl]-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

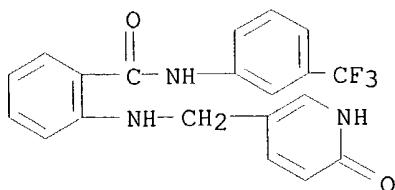
IT 524941-35-3P 524941-36-4P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of anthranilic acid amides and their use as vascular
 endothelial growth factor receptor tyrosine kinase inhibitors)

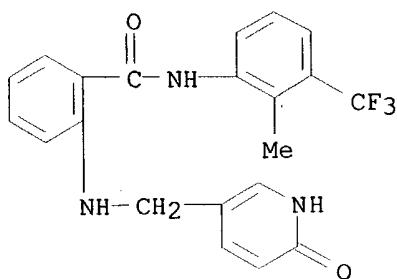
RN 524941-35-3 CAPLUS

CN Benzamide, 2-[(1,6-dihydro-6-oxo-3-pyridinyl)methyl]amino-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



RN 524941-36-4 CAPLUS

CN Benzamide, 2-[(1,6-dihydro-6-oxo-3-pyridinyl)methyl]amino-N-[2-methyl-3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

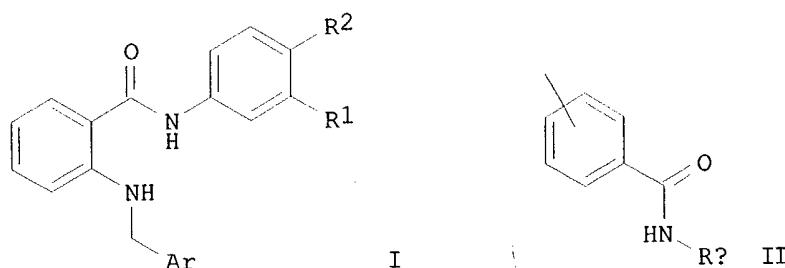
L31 ANSWER 9 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2003:376824 CAPLUS
 DOCUMENT NUMBER: 138:368777
 TITLE: Preparation of pyridyl-substituted anthranilic acid amides for treating neoplastic disease
 INVENTOR(S): Bold, Guido; Furet, Pascal; Manley, Paul William
 PATENT ASSIGNEE(S): Novartis AG, Switz.; Novartis Pharma GmbH
 SOURCE: PCT Int. Appl., 33 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003040101	A1	20030515	WO 2002-EP12445	20021107
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LT, LU, LV, MA, MD, MK, MN, MX, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SE, SG, SI, SK, TJ, TM, TN, TR, TT, UA, US, UZ, VC, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR				
PRIORITY APPLN. INFO.:			GB 2001-26901	A 20011108
			GB 2002-12917	A 20020605

OTHER SOURCE(S): MARPAT 138:368777

ED Entered STN: 16 May 2003

GI



AB The title compds. [I; Ar = II (wherein Ra = H, alkyl; and R1 = H, perfluoroalkyl; R2 = H, halo, alkyl, alkenyl, alkynyl); or Ar = 4-pyridyl

and R1 = perfluoroalkyl; R2 = Br, I, alkyl, alkenyl, alkynyl; or R1 = H, and R2 = F, Br, I, Et, alkyl, alkenyl or alkynyl] and their N-oxides and salts, useful for the treatment esp. of a neoplastic disease, such as a tumor disease, of retinopathy or age-related macular degeneration in the human or animal body, were prep'd. and formulated. Thus, reductive amination of 4-pyridinecarboxaldehyde with 2-amino-N-(4-bromo-3-trifluoromethylphenyl)benzamide (prepn. given) in the presence of NaBH3CN afforded I [Ar = 4-pyridyl; R1 = CF3; R2 = Br]. The IC50-values that can be found for the compds. I are in range of 0.001 to 1 .mu.M in test for activity against VEGF-receptor tyrosine kinase.

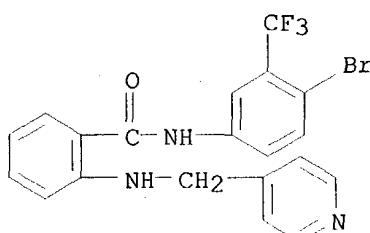
IT 524728-98-1P 524728-99-2P 524729-02-0P

524729-04-2P

RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(prepn. of pyridyl-substituted anthranilic acid amides for treating neoplastic disease)

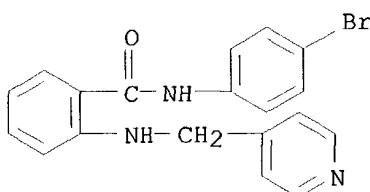
RN 524728-98-1 CAPLUS

CN Benzamide, N-[4-bromo-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



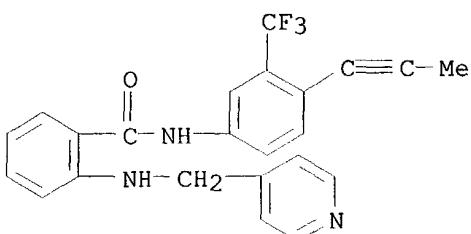
RN 524728-99-2 CAPLUS

CN Benzamide, N-(4-bromophenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 524729-02-0 CAPLUS

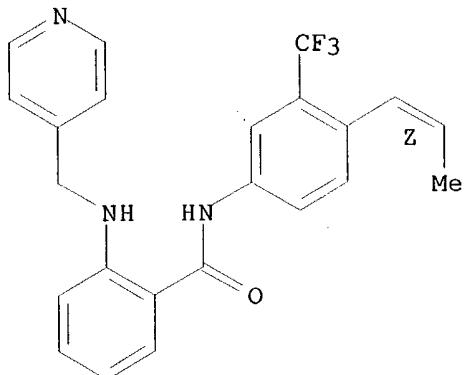
CN Benzamide, N-[4-(1-propynyl)-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 524729-04-2 CAPLUS

CN Benzamide, N-[4-(1Z)-1-propenyl-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]-, monohydrochloride (9CI) (CA INDEX NAME)

Double bond geometry as shown.



● HCl

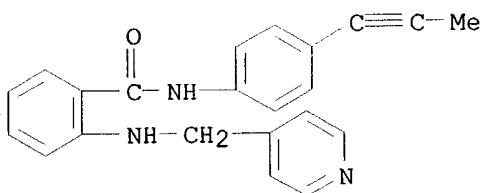
IT 524729-03-1P 524729-05-3P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of pyridyl-substituted anthranilic acid amides for treating neoplastic disease)

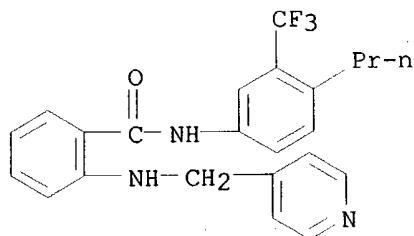
RN 524729-03-1 CAPLUS

CN Benzamide, N-[4-(1-propynyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 524729-05-3 CAPLUS

CN Benzamide, N-[4-propyl-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



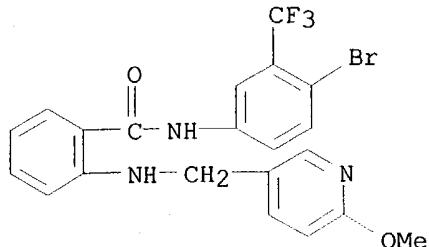
IT 524728-97-0P 524729-01-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of pyridyl-substituted anthranilic acid amides for treating neoplastic disease)

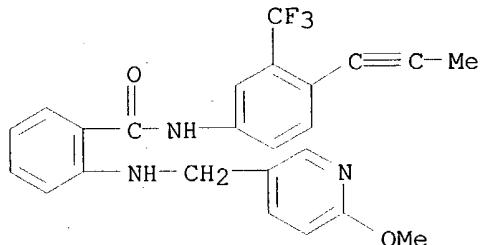
RN 524728-97-0 CAPLUS

CN Benzamide, N-[4-bromo-3-(trifluoromethyl)phenyl]-2-[[[6-methoxy-3-pyridinyl)methyl]amino]- (9CI) (CA INDEX NAME)



RN 524729-01-9 CAPLUS

CN Benzamide, 2-[[[6-methoxy-3-pyridinyl)methyl]amino]-N-[4-(1-propynyl)-3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

~~ANSWER 10 OF 55~~ CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2003:42101 CAPLUS

DOCUMENT NUMBER: 138:106502

TITLE: Preparation of biphenylcarboxylic acid amides as inhibitors of microsomal triglyceride transfer protein (MTP)

INVENTOR(S): Priepke, Henning; Hauel, Norbert; Dahmann, Georg; Thomas, Leo; Mark, Michael

PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma K.-G., Germany

SOURCE: PCT Int. Appl., 193 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

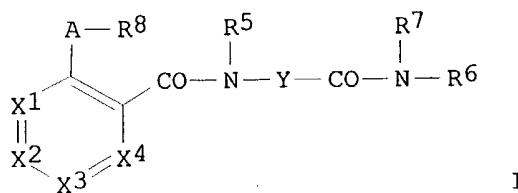
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003004020	A1	20030116	WO 2002-EP7215	20020629
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,				

UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU,
 TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
 CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
 BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
 DE 10132686 A1 20030116 DE 2001-10132686 20010705
 US 2003073836 A1 20030417 US 2002-187860 20020702
 PRIORITY APPLN. INFO.: DE 2001-10132686 A 20010705
 US 2001-304584P P 20010711

OTHER SOURCE(S): MARPAT 138:106502

ED Entered STN: 17 Jan 2003

GI



AB Title compds. I [X1 = CR1; X2 = CR2; X3 = CR3; X4 = CR4; with 1-2 of the groups being a N-atom; R1, R2, R3, R4 = H, halo, alkyl, etc.; A = O, S, NH, etc.; R8 = (un)substituted Ph, 1-naphthyl, 2-naphthyl, etc.; R5 = H, (un)substituted alkyl; R6 = H, alkyl; R7 = (un)substituted alkyl; Y = 1-2 carbon atom(s) bound to (un)substituted 5-membered heteroaryl] and their pharmaceutically acceptable salts were prep'd. For example, coupling of acid II, e.g., prep'd. from 4-hydrazinobenzonitrile in 5-steps, and 4-phenylbenzylamine afforded biphenylcarboxylic acid amide III in 86% yield. In triglyceride transfer protein inhibition studies, compds. I exhibited IC50 values < 100.μM. Compds. I are claimed useful as inhibitors of microsomal triglyceride transfer protein (MTP) for the treatment of atherosclerosis.

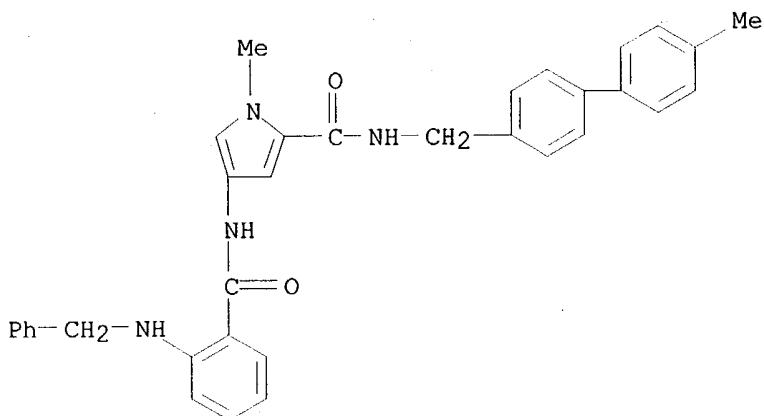
IT 486436-62-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; prepn. of biphenylcarboxylic acid amides as microsomal triglyceride transfer protein (MTP) inhibitors)

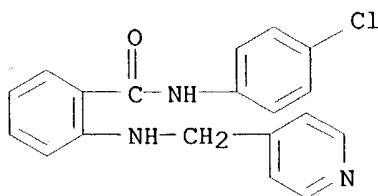
RN 486436-62-8 CAPLUS

CN 1H-Pyrrole-2-carboxamide, 1-methyl-N-[(4'-methyl[1,1'-biphenyl]-4-yl)methyl]-4-[[2-[(phenylmethyl)amino]benzoyl]amino]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

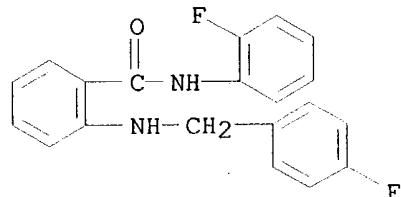
131 ANSWER 11 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2003:665525 CAPLUS
 DOCUMENT NUMBER: 139:345320
 TITLE: Identification of a new chemical class of potent angiogenesis inhibitors based on conformational considerations and database searching
 AUTHOR(S): Furet, Pascal; Bold, Guido; Hofmann, Francesco; Manley, Paul; Meyer, Thomas; Altmann, Karl-Heinz
 CORPORATE SOURCE: Oncology Research, Novartis Pharma AG, Basel, CH-4002, Switz.
 SOURCE: Bioorganic & Medicinal Chemistry Letters (2003), 13(18), 2967-2971
 CODEN: BMCL8; ISSN: 0960-894X
 PUBLISHER: Elsevier Science B.V.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 ED Entered STN: 26 Aug 2003
 AB The vascular endothelial growth factor (VEGF) tyrosine kinase receptors KDR and Flt-1 are targets of current interest in anticancer drug research. PTK787/ZK222584 is a potent inhibitor of these enzymes in clin. evaluation as an antiangiogenic agent. The development of a hypothesis concerning the bioactive conformation of this compd. has led to the discovery of a new class of potent inhibitors of KDR and Flt-1, the anthranilamides. This could be achieved with a limited exptl. effort, which only involved the testing of one archive compd. and the synthesis and testing of one appropriate analog.
 IT 269390-69-4P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (identification, synthesis and structure-activity relationship studies on a new chem. class of potent angiogenesis inhibitors (anthranilamides)-based on conformational considerations and database searching)
 RN 269390-69-4 CAPLUS
 CN Benzamide, N-(4-chlorophenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



IT 618359-41-4

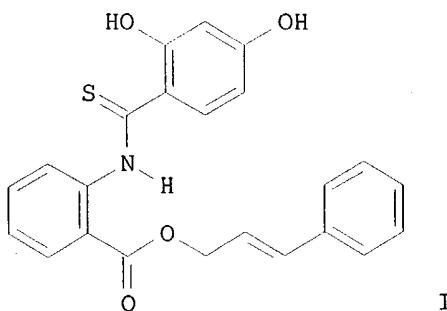
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (identification, synthesis and structure-activity relationship studies on a new chem. class of potent angiogenesis inhibitors (anthranilamides)-based on conformational considerations and database searching)

RN 618359-41-4 CAPLUS

CN Benzamide, N-(2-fluorophenyl)-2-[(4-fluorophenyl)methyl]amino- (9CI)
 (CA INDEX NAME)

REFERENCE COUNT: 22 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

✓
 PS1 ANSWER 12 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2002:940418 CAPLUS
 DOCUMENT NUMBER: 138:169931
 TITLE: Synthesis and Fungistatic Activity of New Groups of 2,4-Dihydroxythiobenzoyl Derivatives against Phytopathogenic Fungi
 AUTHOR(S): Legocki, Jan; Matysiak, Joanna; Niewiadomy, Andrzej; Kostecka, Ma-lgorzata
 CORPORATE SOURCE: IPO, Warsaw, 03-236, Pol.
 SOURCE: Journal of Agricultural and Food Chemistry (2003), 51(2), 362-368
 CODEN: JAFCAU; ISSN: 0021-8561
 PUBLISHER: American Chemical Society
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 138:169931
 ED Entered STN: 12 Dec 2002
 GI



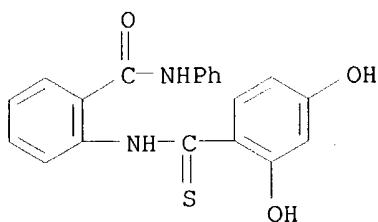
AB Twenty-six compds., derivs. of amides, hydrazines, hydrazides, hydrazone, and semicarbazides, with a 2,4-dihydroxythiobenzoyl moiety, were synthesized from sulfinyl-bis(2,4-dihydroxythiobenzoyl). The compns. and chem. structures of these compds. were confirmed by IR, 1H NMR, EI-MS, and elemental anal. Antifungal properties of chms. under in vitro conditions against five phytopathogenic fungi were estd. In vivo studies against Erysiphe graminis were also carried out. The compds. N-substituted with an 2,4-dihydroxythiobenzamide group proved to be the most active. N-2-(1-Cinnamylbenzene ester)-2,4-dihydroxythiobenzamide (I), under in vitro conditions, showed activity at the level of 80-100% development of most pathogens at a concn. of 20 .mu.g/mL and partially at a concn. of 200 .mu.g/mL. For compds. with -HN-NH- or -NH-N: moiety, weak or no fungistatic properties were found at the concns. studied.

IT 497156-37-3P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn., structure-activity relationship and fungicidal activity of dihydroxythiobenzoyl derivs. via nucleophilic substitution reactions of sulfinylbis(dihydroxythiobenzoyl) with various nucleophiles)

RN 497156-37-3 CAPLUS

CN Benzamide, 2-[(2,4-dihydroxyphenyl)thioxomethyl]amino]-N-phenyl- (9CI)
(CA INDEX NAME)



REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

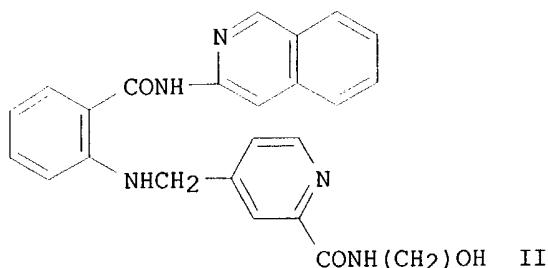
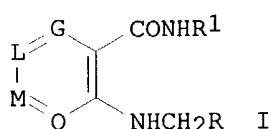
431 ANSWER 13 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:868928 CAPLUS
DOCUMENT NUMBER: 137:352900
TITLE: Selective anthranilamide pyridine amides as inhibitors of VEGFR-2 and VEGFR-3
INVENTOR(S): Ernst, Alexander; Huth, Andreas; Krueger, Martin; Thierauch, Karl-Heinz; Menrad, Andreas; Haberey, Martin
PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany
SOURCE: PCT Int. Appl., 115 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002090352	A2	20021114	WO 2002-EP4924	20020503
WO 2002090352	A3	20030501		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
DE 10123574	A1	20021128	DE 2001-10123574	20010508
DE 10125294	A1	20021121	DE 2001-10125294	20010515
DE 10164590	A1	20030710	DE 2001-10164590	20011221
EP 1392680	A2	20040303	EP 2002-735333	20020503
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
PRIORITY APPLN. INFO.:			DE 2001-10123574 A	20010508
			DE 2001-10125294 A	20010515
			DE 2001-10164590 A	20011221
			WO 2002-EP4924 W	20020503

OTHER SOURCE(S): MARPAT 137:352900

ED Entered STN: 15 Nov 2002
GI

AB Title compds. I [G, L, M, Q = N, (un)substituted CH, .1toreq.1 of them being N; R = (un)substituted N heterocycle; R1 = (un)substituted alkyl, alkenyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl] were prep'd. I are inhibitors of VEGFR-2 and VEGFR-3 and are used as medicaments for treating diseases that are caused by persistent angiogenesis, such as psoriasis, Kaposi's sarcoma, restenosis, such as e.g. stent-induced restenosis, endometriosis, Crohn's disease, Hodgkin's disease, leukemia, arthritis, such as rheumatoid arthritis, hemangioma, angiofibromatosis, in eye diseases such as diabetic retinopathy, neovascular glaucoma, in kidney diseases such as glomerulonephritis, diabetic nephropathy, malign nephrosclerosis, thrombic micro-angiopathic syndrome, transplant rejection and glomerulopathy, in fibrotic diseases such as hepatic cirrhosis, mesangial-cell proliferative diseases, arteriosclerosis, damage to the nerve tissue and inhibition of the re-occlusion of vessels after balloon

catheter treatment, in vessel prosthetics or after the use of mech. devices for keeping vessels open, e.g. stents, as immunosuppressants, to support wound healing without scars and in cases of age spots and contact dermatitis. I can also be used as inhibitors of VEGFR-3 in lymphangiogenesis for hyperplastic and dysplastic changes in the lymphatic system. Thus, 2-amino-N-isoquinolin-3-ylbenzamide was treated with 2-bromo-5-pyridinecarboxaldehyde, followed by carboxylation and amidation to give the amide II. II had IC₅₀ for inhibition of VEGFR-2 of 40 nM and for inhibition of cytochrome 450 isoenzyme 2C9 of 2.9 .mu.M.

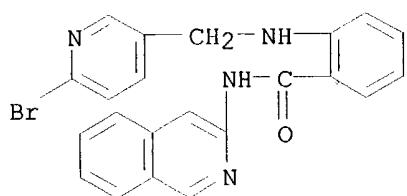
IT 474799-36-5P 474799-37-6P 474799-38-7P

474799-46-7P 474799-57-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of isoquinolinylcarbamoylphenylaminomethylpyridinecarboxamides as VEGFR-2 and VEGFR-3 inhibitors)

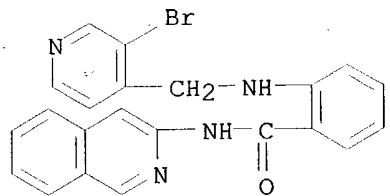
RN 474799-36-5 CAPLUS

CN Benzamide, 2-[[6-bromo-3-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI)
(CA INDEX NAME)



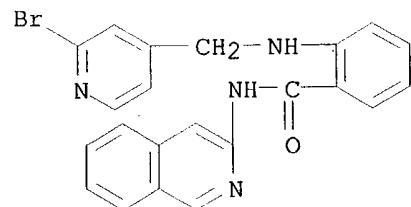
RN 474799-37-6 CAPLUS

CN Benzamide, 2-[[3-bromo-4-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI)
(CA INDEX NAME)



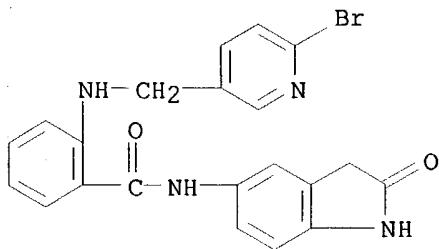
RN 474799-38-7 CAPLUS

CN Benzamide, 2-[[2-bromo-4-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI)
(CA INDEX NAME)

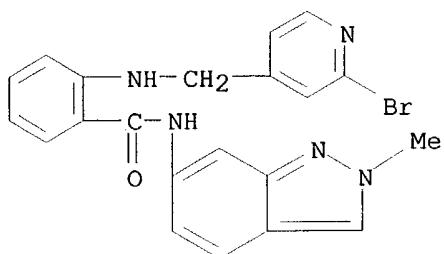


RN 474799-46-7 CAPLUS

CN Benzamide, 2-[[6-bromo-3-pyridinyl)methyl]amino]-N-(2,3-dihydro-2-oxo-1H-indol-5-yl)- (9CI) (CA INDEX NAME)



RN 474799-57-0 CAPLUS
 CN Benzamide, 2-[[[(2-bromo-4-pyridinyl)methyl]amino]-N-(2-methyl-2H-indazol-6-yl)-(9CI) (CA INDEX NAME)

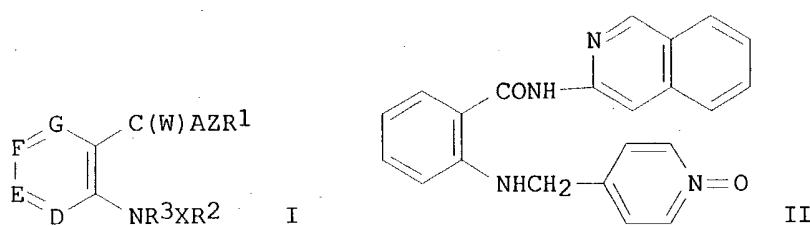


L31 ANSWER 14 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2002:868925 CAPLUS
 DOCUMENT NUMBER: 137:352899
 TITLE: Pyridylmethylanthranilamide N-oxides as inhibitors of VEGFR II kinase
 INVENTOR(S): Ernst, Alexander; Huth, Andreas; Krueger, Martin; Thierauch, Karl-Heinz; Menrad, Andreas; Haberey, Martin
 PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany
 SOURCE: PCT Int. Appl., 50 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002090349	A1	20021114	WO 2002-EP4923	20020503
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
DE 10123573	A1	20021128	DE 2001-10123573	20010508
DE 10125293	A1	20021121	DE 2001-10125293	20010515
EP 1389201	A1	20040218	EP 2002-740563	20020503
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
PRIORITY APPLN. INFO.:			DE 2001-10123573 A	20010508
			DE 2001-10125293 A	20010515

WO 2002-EP4923 W 20020503

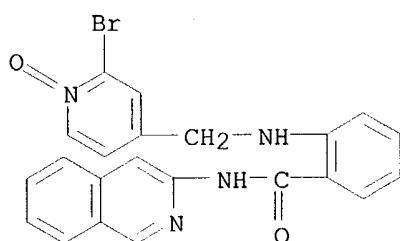
OTHER SOURCE(S): MARPAT 137:352899
ED Entered STN: 15 Nov 2002 ..
GI



AB Title compds. I [D, E, F, G = N, (un)substituted CH; A = (un)substituted NH; W = O, S, H2, (un)substituted NH; X, Z = (un)substituted alkylene; R1 = (un)substituted alkyl, alkenyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; R2 = (un)substituted hetaryl N-oxide; R3 = H, alkyl] were prepd. These compds. can be used in the treatment of psoriasis, Kaposi's sarcoma, restenosis, endometriosis, Crohn's disease, Hodgkin's disease, leukemia, arthritis, such as rheumatoid arthritis, hemangioma, angiofibroma, eye diseases, such as diabetic retinopathy, neovascular glaucoma, renal diseases, such as glomerulonephritis, diabetic nephropathy, malignant nephrosclerosis, thrombotic microangiopathic syndromes, transplant rejections and glomerulopathy, fibrotic diseases such as cirrhosis of the liver, mesangial cell-proliferative diseases, arteriosclerosis, injuries of the nerve tissue, and for inhibiting the reocclusion of vessels after balloon catheter treatment, for use in vascular prosthetics or after inserting mech. devices for holding vessels open such as, e.g. stents, as immunosuppressants, as an aid in scar-free wound healing, and for treating age spots and contact dermatitis. They can also be used as VEGFR-3 inhibitors in lymphangiogenesis. Thus, the N-oxide II was obtained by reductive alkylation of 2-amino-N-isoquinolin-3-ylbenzamide with isonicotinaldehyde N-oxide and had IC50 for inhibition of VEGFR II of 0.03 .mu.M.

IT 474760-12-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of pyridylmethylanthranilamide N-oxides as inhibitors of VEGFR)

RN 474760-12-8 CAPLUS
CN Benzamide, 2-[(2-bromo-1-oxido-4-pyridinyl)methyl]amino]-N-3-isquinolinalyl- (9CI) (CA INDEX NAME)

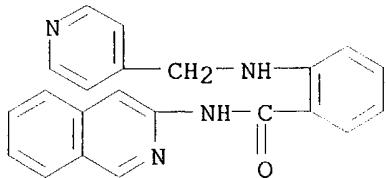


IT 267891-44-1

RL: RCT (Reactant); RACT (Reactant or reagent)
 (prepn. of pyridylmethylanthranilamide N-oxides as inhibitors of VEGFR
 II kinase)

RN 267891-44-1 CAPLUS

CN Benzamide, N-3-isoquinolinyl-2-[(4-pyridinylmethyl)amino]- (9CI) (CA
 INDEX NAME)



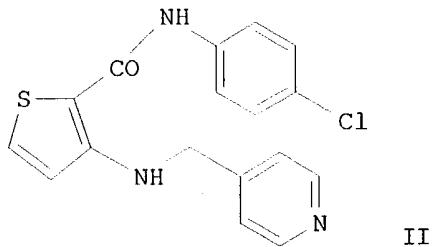
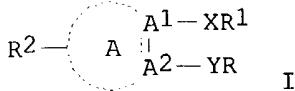
REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L31 ANSWER 15 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2002:658116 CAPLUS
 DOCUMENT NUMBER: 137:201332
 TITLE: Preparation of heterocyclalkylamine derivatives as
 remedies for angiogenesis mediated diseases
 INVENTOR(S): Chen, Guoqing; Adams, Jeffrey; Bemis, Jean; Booker,
 Shon; Cai, Guolin; Croghan, Michael; Dipietro, Lucian;
 Dominguez, Celia; Elbaum, Daniel; Germain, Julie;
 Geuns-meyer, Stephanie; Handley, Michael; Huang, Qi;
 Kim, Joseph L.; Kim, Tae-seong; Kiselyov, Alexander;
 Ouyang, Xiaohu; Patel, Vinod F.; Smith, Leon M.; Stec,
 Markian; Tasker, Andrew; Xi, Ning; Xu, Shimin; Yuan,
 Chester Chenguang
 PATENT ASSIGNEE(S): Amgen Inc., USA
 SOURCE: PCT Int. Appl., 502 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002066470	A1	20020829	WO 2002-US743	20020111
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2003125339	A1	20030703	US 2002-46681	20020110
BR 2002006435	A	20030923	BR 2002-6435	20020111
EP 1358184	A1	20031105	EP 2002-717325	20020111
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
EE 200300324	A	20031215	EE 2003-324	20020111
NO 2003003181	A	20030911	NO 2003-3181	20030711
PRIORITY APPLN. INFO.:			US 2001-261339P	P 20010112
			US 2001-323764P	P 20010919
			US 2002-46681	A 20020110

WO 2002-US743 W 20020111

OTHER SOURCE(S): MARPAT 137:201332
 ED Entered STN: 30 Aug 2002
 GI

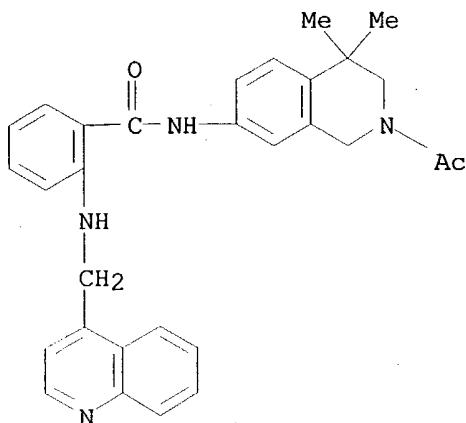


AB Title compds. [I; A1, A2 independently = C, N; A = 5-, or 6-membered partially satd. heterocyclyl, 5-, or 6-membered heterocyclyl, 9-, or 10-membered fused partially satd. heterocyclyl, 9-, 10-, or 11-membered fused heteroaryl, naphthyl, 4-, 5-, or 6-membered cycloalkenyl; X = C:ZNR3, C:ZN(R3)R4; Z = O, S; Y = N:CH, NR5(CR6R7), R8N(R5)(CR6R7), NR5(CR6R7)R8; R = 5-, or 6-membered (un)substituted heterocyclyl, 9-, 10-, 11-membered heterocyclyl; R1 = 6-10-membered (un)substituted aryl, 5-, or 6-membered (un)substituted heterocyclyl, 9-11 membered (un)substituted fused heterocyclyl, cycloalkyl, cycloalkenyl; R2 = H, halo, oxo, SH, COOH, CHO; R3 = H, alkyl, 5-, or 6-membered heterocyclyl; R4 = alkylene, alkenylene, alkynylene; R5 = H, alkyl, aralkyl, C6H5; R6, R7 independently = H, halo, CN, alkyl; R6R7 = cycloalkyl; R8 = alkylene, etc.] are prepd. and are effective for prophylaxis and treatment of diseases, such as angiogenesis mediated diseases. The invention encompasses novel compds., analogs, prodrugs and pharmaceutically acceptable derivs. thereof, pharmaceutical compns. and methods for prophylaxis and treatment of diseases and other maladies or conditions involving, cancer and the like. The subject invention also relates to processes for making such compds. as well as to intermediates useful in such processes. Thus, the title compd. II was prepd. from Me 3-amino-2-thiophenecarboxylate, 4-chloroaniline, and 4-pyridine carboxaldehyde via coupling reaction.

IT **453564-10-8P**
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of heterocyclylalkylamine derivs. as remedies for angiogenesis mediated diseases)

RN 453564-10-8 CAPLUS

CN Benzamide, N-(2-acetyl-1,2,3,4-tetrahydro-4,4-dimethyl-7-isoquinolinyl)-2-[(4-quinolinylmethyl)amino]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

~~131~~ ANSWER 16 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:171866 CAPLUS

DOCUMENT NUMBER: 136:232313

TITLE: Preparation of pyrimidine derivatives as G protein-coupled receptor kinase (GRK) inhibitors

INVENTOR(S): Fukumoto, Shoji; Watanabe, Toshifumi; Ikeda, Shota

PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan

SOURCE: PCT Int. Appl., 322 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

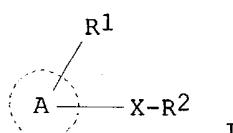
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002018350	A1	20020307	WO 2001-JP7397	20010829
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2001082520	A5	20020313	AU 2001-82520	20010829
JP 2002145778	A2	20020522	JP 2001-259683	20010829
PRIORITY APPLN. INFO.:			JP 2000-264499	A 20000829
			WO 2001-JP7397	W 20010829

OTHER SOURCE(S): MARPAT 136:232313

ED Entered STN: 08 Mar 2002

GI



AB Disclosed are novel GRK inhibitors which contains compds. represented by the formula (I), a salt thereof, or a prodrug comprising either of these (wherein ring A represents optionally further substituted nitrogen-contg. heterocycle; R1 and R2 each represents optionally substituted amino; and X represents a spacer comprising a linear part constituted of one to four atoms, provided that R1 may be bonded to R2 or/and X to form a ring). They are useful as preventives/remedies for cardiac failure. Thus, 5.48 g K₂CO₃ and 7.52 g 2-aminophenyl 2-nitrophenyl sulfide were added to a suspension of 5.61 g 4-amino-5-bromomethyl-2-methylpyrimidine hydrobromide in 40 mL acetone at room temp. and stirred at 65.degree. for 64 h to give 2.36 g N-[(4-amino-2-methyl-5-pyrimidinyl)methyl]-N-[2-[(2-nitrophenyl)thio]phenyl]amine (II). All 10 compds. tested including II at 30 .mu.M inhibited 30% human GRK2 expressed by human GRK2 gene in COS-7 cells. A capsule and a tablet formulation contg. II were also prepd.

IT 403515-67-3P 403515-68-4P 403515-69-5P

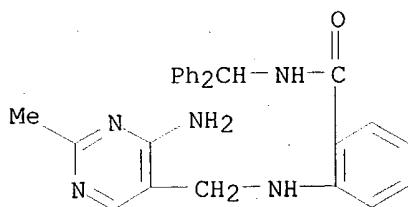
403515-71-9P 403515-72-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of pyrimidine derivs. as G protein-coupled receptor kinase (GRK) inhibitors for prevention and/or treatment for cardiac failure)

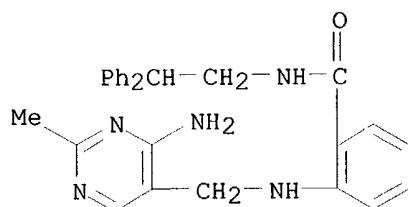
RN 403515-67-3 CAPLUS

CN Benzamide, 2-[[[(4-amino-2-methyl-5-pyrimidinyl)methyl]amino]-N-(diphenylmethyl)- (9CI) (CA INDEX NAME)



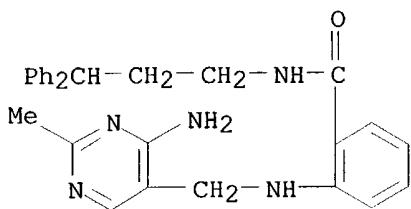
RN 403515-68-4 CAPLUS

CN Benzamide, 2-[[[(4-amino-2-methyl-5-pyrimidinyl)methyl]amino]-N-(2,2-diphenylethyl)- (9CI) (CA INDEX NAME)



RN 403515-69-5 CAPLUS

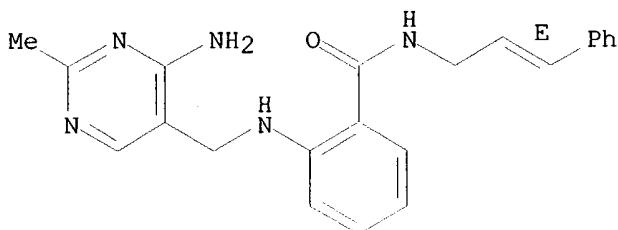
CN Benzamide, 2-[[[(4-amino-2-methyl-5-pyrimidinyl)methyl]amino]-N-(3,3-diphenylpropyl)- (9CI) (CA INDEX NAME)



RN 403515-71-9 CAPLUS

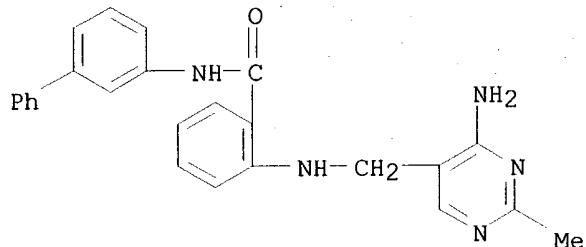
CN Benzamide, 2-[[(4-amino-2-methyl-5-pyrimidinyl)methyl]amino]-N-[(2E)-3-phenyl-2-propenyl]- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



RN 403515-72-0 CAPLUS

CN Benzamide, 2-[[(4-amino-2-methyl-5-pyrimidinyl)methyl]amino]-N-[1,1'-biphenyl]-3-yl- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 17 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:882097 CAPLUS

DOCUMENT NUMBER: 137:384763

TITLE: Preparation of cyanoanthranylamides as vascular endothelial growth factor (VEGF) receptor inhibitors

INVENTOR(S): Huth, Andreas; Krueger, Martin; Thierauch, Karl-Heinz; Ernst, Alexander; Menrad, Andreas; Haberey, Martin

PATENT ASSIGNEE(S): Schering Ag, Germany

SOURCE: Ger. Offen., 14 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2

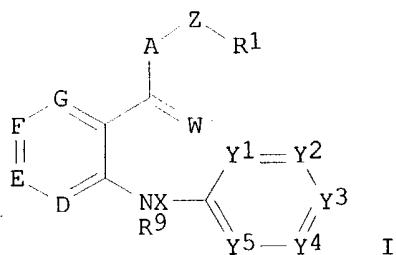
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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DE 10125295 A1 20021121 DE 2001-10125295 20010515
 WO 2003000678 A1 20030103 WO 2002-EP4921 20020503
 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
 CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM,
 HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
 LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL,
 PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA,
 UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
 CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
 BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
 EP 1387838 A1 20040211 EP 2002-748691 20020503
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
 PRIORITY APPLN. INFO.: DE 2001-10123587 A 20010508
 DE 2001-10125295 A 20010515
 WO 2002-EP4921 W 20020503

OTHER SOURCE(S): MARPAT 137:384763

ED Entered STN: 21 Nov 2002
 GI



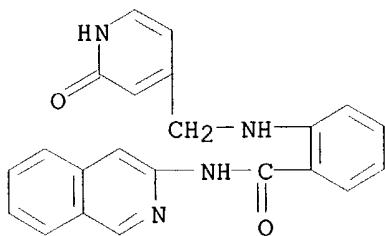
AB Title compds. [I; A = NR7; W = O, S, 2H, NR8; Z = bond, NR10, :N, (branched) (substituted) alkyl; X = alkyl; R1 = (substituted) (branched) alkyl, alkenyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; Y1-Y5 = N, CY6; Y6 = cyano, halo, alkyl, alkoxy, amino, OH (with the proviso that the ring contains at least one of N and is substituted with at least one of cyano group); D = N, CR3; E = N, CR4; F = N, CR5; G = N, CR6; R3-R6 = H, halo, (substituted) alkoxy, alkyl, carboxyalkyl; R7 = H, alkyl, etc.; R8-R10 = H, alkyl], were prep'd. Thus, N-(isoquinolin-3-yl)-2-(4-pyridylmethyl)aminobenzoic acid amide N-oxide was treated one after another with DMF, Et3N, and Me3SiCN followed by heating the bath temp. at 110.degree. to give 14% N-(isoquinolin-3-yl)-2-[4-(2-cyanopyridyl)methyl]aminobenzoic acid amide. The latter inhibited the tyrosine kinase receptor VEGFR II (KDR) with IC50 = 1.times.10⁻⁸ mM.

IT 267891-90-7

RL: RCT (Reactant); RACT (Reactant or reagent)
 (prepn. of cyanoanthranylamides as vascular endothelial growth factor (VEGF) receptor inhibitors)

RN 267891-90-7 CAPLUS

CN Benzamide, 2-[[1,2-dihydro-2-oxo-4-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI) (CA INDEX NAME)



ANSWER 18 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:882056 CAPLUS

DOCUMENT NUMBER: 137:384762

TITLE: Preparation of cyanoanthranyl amides as vascular endothelial growth factor (VEGF) receptor inhibitors

INVENTOR(S): Huth, Andreas; Krueger, Martin; Ernst, Alexander; Thierauch, Karl-Heinz; Haberey, Martin; Menrad, Andreas

PATENT ASSIGNEE(S): Schering Ag, Germany

SOURCE: Ger. Offen., 14 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2

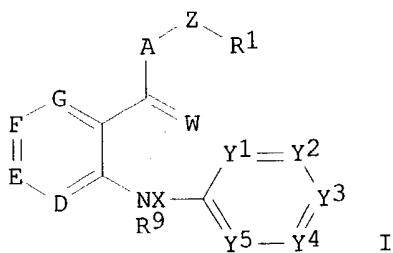
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10123587	A1	20021121	DE 2001-10123587	20010508
WO 2003000678	A1	20030103	WO 2002-EP4921	20020503
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1387838	A1	20040211	EP 2002-748691	20020503
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
PRIORITY APPLN. INFO.:			DE 2001-10123587 A	20010508
			DE 2001-10125295 A	20010515
			WO 2002-EP4921 W	20020503

OTHER SOURCE(S): MARPAT 137:384762

ED Entered STN: 21 Nov 2002

GI



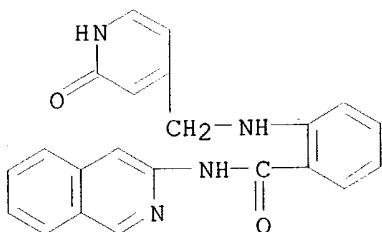
AB Title compds. [I; A = NR7; W = O, S, 2H, NR8; Z = bond, NR10, :N, (branched) (substituted) alkyl; X = alkyl; R1 = (substituted) (branched) alkyl, alkenyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; Y1-Y5 = N, CY6; Y6 = cyano, halo, alkyl, alkoxy, amino, OH (with the proviso that the ring contains at least one of N and is substituted with at least one of cyano group); D = N, CR3; E = N, CR4; F = N, CR5; G = N, CR6; R3-R6 = H, halo, (substituted) alkoxy, alkyl, carboxyalkyl; R7 = H, alkyl, etc.; R8-R10 = H, alkyl], were prep'd. Thus, N-(isoquinolin-3-yl)-2-(4-pyridylmethyl)aminobenzoic acid amide N-oxide was treated one after another with DMF, Et3N, and Me3SiCN followed by heating the bath temp. at 110.degree. to give 14% N-(isoquinolin-3-yl)-2-[4-(2-cyanopyridyl)methyl]aminobenzoic acid amide. The latter inhibited the tyrosine kinase receptor VEGFR II (KDR) with IC50 = 1.times.10⁻⁸ mM.

IT 267891-90-7

RL: RCT (Reactant); RACT (Reactant or reagent)
(prepn. of cyanoanthranylamides as vascular endothelial growth factor (VEGF) receptor inhibitors)

RN 267891-90-7 CAPLUS

CN Benzamide, 2-[[[(1,2-dihydro-2-oxo-4-pyridinyl)methyl]amino]-N-3-isoquinolinyl- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ESI ANSWER 19 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:880425 CAPLUS

DOCUMENT NUMBER: 138:106488

TITLE: Anthranilic Acid Amides: A Novel Class of Antiangiogenic VEGF Receptor Kinase Inhibitors

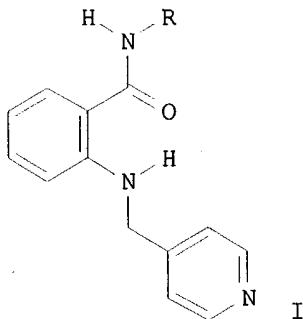
AUTHOR(S): Manley, Paul W.; Furet, Pascal; Bold, Guido; Brueggen, Josef; Mestan, Juergen; Meyer, Thomas; Schnell, Christian R.; Wood, Jeanette; Haberey, Martin; Huth, Andreas; Krueger, Martin; Menrad, Andreas; Ottow, Eckhard; Seidelmann, Dieter; Siemeister, Gerhard; Thierauch, Karl-Heinz

CORPORATE SOURCE: Oncology Research, Novartis Pharma AG, Basel, CH-4057, Switz.

SOURCE: Journal of Medicinal Chemistry (2002), 45(26),

5687-5693
CODEN: JMCMAR; ISSN: 0022-2623

PUBLISHER: American Chemical Society
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 138:106488
ED Entered STN: 21 Nov 2002
GI



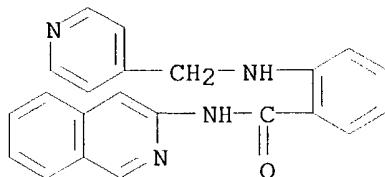
AB Two readily synthesized anthranilamide, VEGF receptor tyrosine kinase inhibitors have been prep'd. and evaluated as angiogenesis inhibitors. 2-[(4-Pyridyl)methyl]amino-N-[3-(trifluoromethyl)phenyl]benzamide [I; R = 3-CF₃C₆H₄ (II)] and N-3-isoquinolinyl-2-[(4-pyridinylmethyl)amino]benzamide [I; R = 3-isoquinolinyl (III)] potently and selectively inhibit recombinant VEGFR-2 and VEGFR-3 kinases. As a consequence of their physicochem. properties, these anthranilamides readily penetrate cells and are absorbed following once daily oral administration to mice. Both II and III potently inhibit VEGF-induced angiogenesis in an implant model, with ED₅₀ values of 7 mg/kg. In a mouse orthotopic model of melanoma, II and III potently inhibited both the growth of the primary tumor as well as the formation of spontaneous peripheral metastases. The anthranilamides II and III represent a new structural class of VEGFR kinase inhibitors, which possess potent antiangiogenic and antitumor properties.

IT 267891-44-1P 269390-77-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(prepn. and antiangiogenic and antitumor activity of VEGF receptor kinase inhibitor anthranilic acid amides)

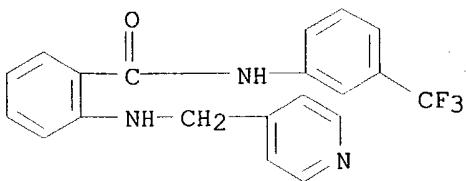
RN 267891-44-1 CAPLUS

CN Benzamide, N-3-isoquinolinyl-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



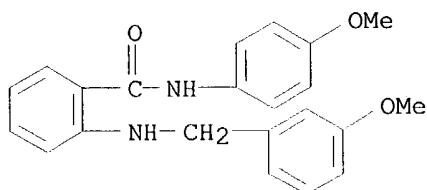
RN 269390-77-4 CAPLUS

CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L31 ANSWER 20 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2002:603273 CAPLUS
 DOCUMENT NUMBER: 138:122629
 TITLE: Synthesis of 1,4-benzodiazepine-2,5-dione derivatives
 AUTHOR(S): Ho, Tong-Ing; Chen, Wen-Shiong; Hsu, Chi-Wei; Tsai, Yeun-Min; Fang, Jim-Min
 CORPORATE SOURCE: Dep. of Chem., National Taiwan Univ., Taipei, Taiwan
 SOURCE: Heterocycles (2002), 57(8), 1501-1506
 CODEN: HTCYAM; ISSN: 0385-5414
 PUBLISHER: Japan Institute of Heterocyclic Chemistry
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 138:122629
 ED Entered STN: 13 Aug 2002
 AB A synthesis of a series of 1,4-benzodiazepine-2,5-dione derivs. with a carboxy group at the 3-position is realized in good yields by using Me malonylchloride as a key reagent and intramol. nucleophilic substitution as ring closure reaction. The synthesis of 4-(4-Methoxyphenyl)-1-[(3-methoxyphenyl)methyl]-2,5-dioxo-1H-1,4-benzodiazepine-3-carboxylic acid Me ester was described.
 IT 489446-50-6P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (prepn. of 2,5-dioxo-1H-1,4-benzodiazepine-3-carboxylate derivs.)
 RN 489446-50-6 CAPLUS
 CN Benzamide, N-(4-methoxyphenyl)-2-[(3-methoxyphenyl)methyl]amino]- (9CI)
 (CA INDEX NAME)



REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

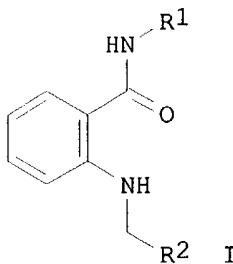
L31 ANSWER 21 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2001:833307 CAPLUS
 DOCUMENT NUMBER: 136:53680
 TITLE: Preparation of anthranilic acid arylamides as inhibitors of tyrosine kinase KDR and FLT.
 INVENTOR(S): Krueger, Martin; Huth, Andreas; Petrov, Orlin; Seidelmann, Dieter; Thierauch, Karl-Heinz; Haberey, Martin; Menrad, Andreas; Ernst, Alexander
 PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany
 SOURCE: PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001085719	A1	20011115	WO 2001-EP5214	20010508
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
DE 10023486	C1	20020314	DE 2000-10023486	20000509
EP 1280799	A1	20030205	EP 2001-940416	20010508
EP 1280799	B1	20040121		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
BR 2001010621	A	20030325	BR 2001-10621	20010508
JP 2003532725	T2	20031105	JP 2001-582320	20010508
AT 258174	E	20040215	AT 2001-1940416	20010508
NO 2002005358	A	20021108	NO 2002-5358	20021108
BG 107261	A	20030630	BG 2002-107261	20021108
PRIORITY APPLN. INFO.:			DE 2000-10023486 A	20000509
			WO 2001-EP5214	W 20010508

OTHER SOURCE(S): MARPAT 136:53680

ED Entered STN: 16 Nov 2001
GI

AB Title compds. [I; R₁ = (substituted) oxobenzopyranyl, quinolinyl, Ph, isoquinolinyl, benzimidazolyl, etc.; R₂ = pyridyl, 2-oxopyridyl, 2-hydroxypyridyl; R₃ = H, F], were prep'd. Thus, N-(2-oxo-2H-1-benzopyran-3-yl)-2-aminobenzamide (prepn. given) was stirred with 4-pyridinecarboxaldehyde in AcOH/MeOH; NaBH₃CN was added to give N-(2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridyl)methyl]aminobenzamide. The latter inhibited KDR with IC₅₀ = 0.003 .mu.M.

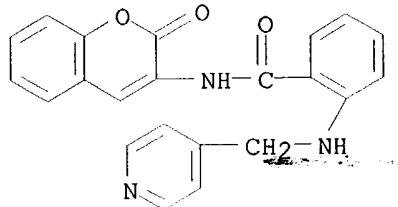
IT 381694-53-7P 381694-55-9P 381694-58-2P
 381694-61-7P 381694-64-0P 381694-67-3P
 381694-70-8P 381694-73-1P 381694-76-4P
 381694-79-7P 381694-82-2P 381694-85-5P
 381694-88-8P 381694-91-3P 381694-94-6P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU

(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of anthranilic acid arylamides as inhibitors of tyrosine kinase KDR and FLT)

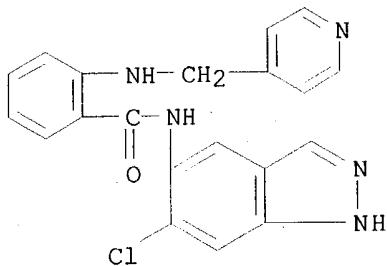
RN 381694-53-7 CAPLUS

CN Benzamide, N-(2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



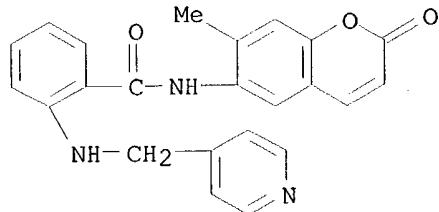
RN 381694-55-9 CAPLUS

CN Benzamide, N-(6-chloro-1H-indazol-5-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



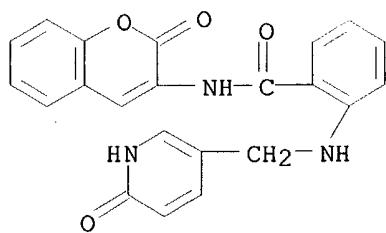
RN 381694-58-2 CAPLUS

CN Benzamide, N-(7-methyl-2-oxo-2H-1-benzopyran-6-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

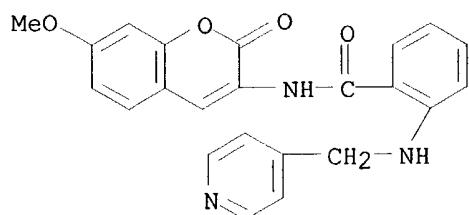


RN 381694-61-7 CAPLUS

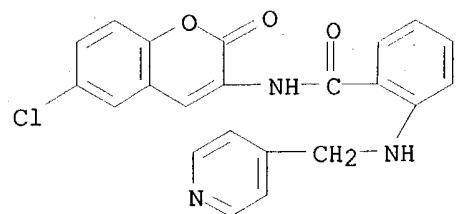
CN Benzamide, 2-[[[(1,6-dihydro-6-oxo-3-pyridinyl)methyl]amino]-N-(2-oxo-2H-1-benzopyran-3-yl)- (9CI) (CA INDEX NAME)



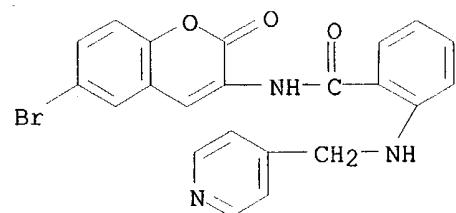
RN 381694-64-0 CAPLUS
CN Benzamide, N-(7-methoxy-2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



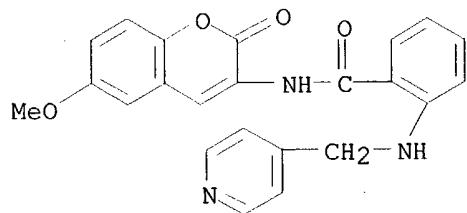
RN 381694-67-3 CAPLUS
CN Benzamide, N-(6-chloro-2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



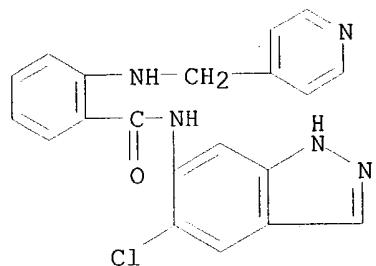
RN 381694-70-8 CAPLUS
CN Benzamide, N-(6-bromo-2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



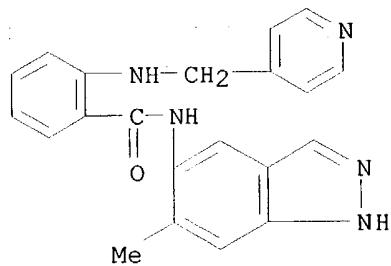
RN 381694-73-1 CAPLUS
CN Benzamide, N-(6-methoxy-2-oxo-2H-1-benzopyran-3-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



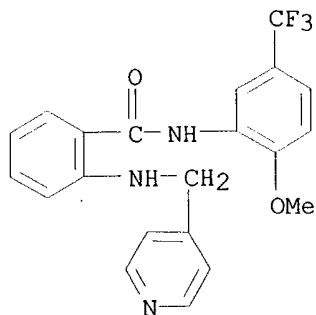
RN 381694-76-4 CAPLUS
CN Benzamide, N-(5-chloro-1H-indazol-6-yl)-2-[(4-pyridinylmethyl)amino]-
(9CI) (CA INDEX NAME)



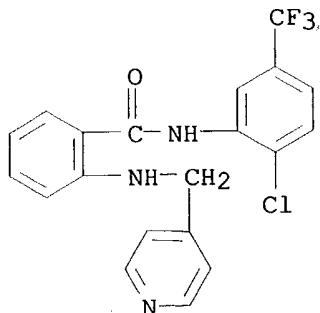
RN 381694-79-7 CAPLUS
CN Benzamide, N-(6-methyl-1H-indazol-5-yl)-2-[(4-pyridinylmethyl)amino]-
(9CI) (CA INDEX NAME)



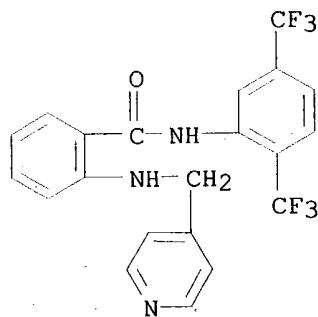
RN 381694-82-2 CAPLUS
CN Benzamide, N-[2-methoxy-5-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



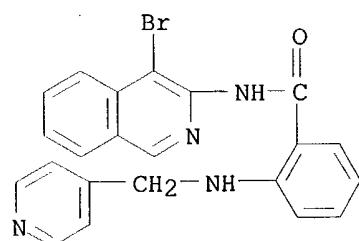
RN 381694-85-5 CAPLUS
CN Benzamide, N-[2-chloro-5-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



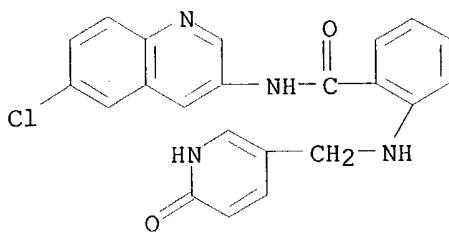
RN 381694-88-8 CAPLUS
CN Benzamide, N-[2,5-bis(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 381694-91-3 CAPLUS
CN Benzamide, N-(4-bromo-3-isoquinolinyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

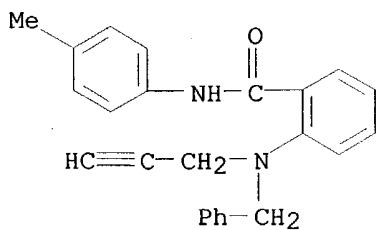


RN 381694-94-6 CAPLUS
CN Benzamide, N-(6-chloro-3-quinolinyl)-2-[(1,6-dihydro-6-oxo-3-pyridinyl)methylamino]- (9CI) (CA INDEX NAME)

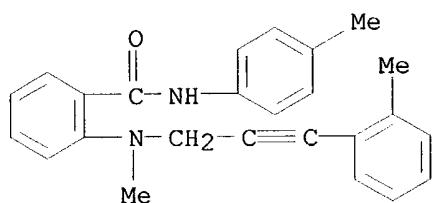


REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

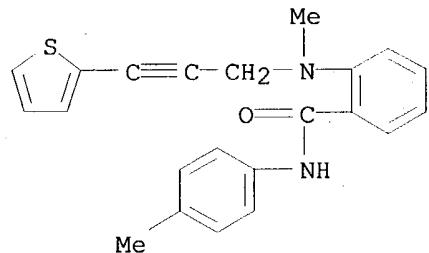
L81 ANSWER 22 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2001:538833 CAPLUS
 DOCUMENT NUMBER: 135:344437
 TITLE: Copper-catalyzed heteroannulation with alkynes: a general and highly regio- and stereoselective method for the synthesis of (E)-2-(2-arylvinyl)quinazolinones
 AUTHOR(S): Kundu, N. G.; Chaudhuri, G.
 CORPORATE SOURCE: Department of Organic Chemistry, Indian Association for Cultivation of Science, Jadavpur, Calcutta, 700 032, India
 SOURCE: Tetrahedron (2001), 57(31), 6833-6842
 CODEN: TETRAB; ISSN: 0040-4020
 PUBLISHER: Elsevier Science Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 ED Entered STN: 26 Jul 2001
 AB A highly regio- and stereoselective procedure for the synthesis of 2-substituted-1,2,3,4-tetrahydroquinazolinones through a two-step procedure, e.g. (i) palladium-copper catalyzed C-arylation of terminal alkynes and (ii) copper-catalyzed cyclization of disubstituted alkynes, is described. 2-[Alkyl(2-propynyl)amino]-N-(4-methylphenyl)benzamides reacted with aryl iodides in the presence of (Ph₃P)₂PdCl₂ (2.5 mol%), CuI (5 mol%), Et₃N (5 equiv.) in CH₃CN at rt for 16 h to yield disubstituted alkynes which could then be cyclized with CuI (20 mol%), K₂CO₃ (2.5 equiv.), Bu₄NBr (1 equiv.) in CH₃CN at 80. degree.C for 16-24 h to yield 1-methyl(benzyl)-(E)-2-(2-arylvinyl)-3-p-tolyl-1,2,3,4-tetrahydro-4-quinazolinones in good yields. Said substituted [[(aminocarbonyl)phenyl]amino]alkynes included N-(4-methylphenyl)-2-[methyl(3-aryl-2-propynyl)amino]benzamide and N-(4-methylphenyl)-2-[(phenylmethyl)(3-aryl-2-propynyl)amino]benzamide derivs. Only in a few cases, benzodiazepinones were obtained in poor yield. The synthesis of novel uracil derivs. was also described.
 IT 350603-00-8P 350603-01-9P 350603-02-0P
 350603-03-1P 371258-55-8P 371258-56-9P
 371258-57-0P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (regioselective, stereoselective prepn. of (E)-2-(2-arylvinyl)quinazolinones via copper-catalyzed heteroannulation of [[(aryl)propynyl]amino]benzamide derivs.)
 RN 350603-00-8 CAPLUS
 CN Benzamide, N-(4-methylphenyl)-2-[(phenylmethyl)-2-propynylamino]- (9CI)
 (CA INDEX NAME)



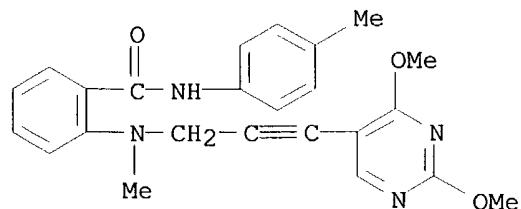
RN 350603-01-9 CAPLUS
 CN Benzamide, 2-[methyl[3-(2-methylphenyl)-2-propynyl]amino]-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



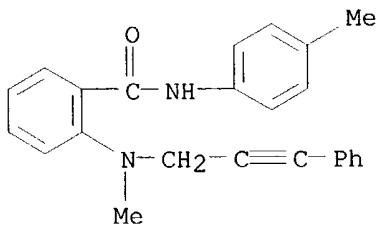
RN 350603-02-0 CAPLUS
 CN Benzamide, N-(4-methylphenyl)-2-[methyl[3-(2-thienyl)-2-propynyl]amino]- (9CI) (CA INDEX NAME)



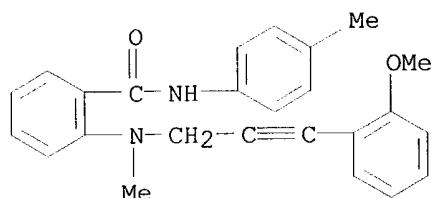
RN 350603-03-1 CAPLUS
 CN Benzamide, 2-[{3-[2,4-dimethoxy-5-pyrimidinyl]-2-propynyl}methylamino]-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



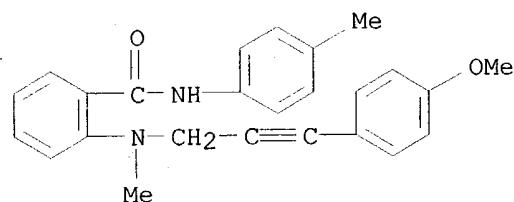
RN 371258-55-8 CAPLUS
 CN Benzamide, N-(4-methylphenyl)-2-[methyl(3-phenyl-2-propynyl)amino]- (9CI) (CA INDEX NAME)



RN 371258-56-9 CAPLUS
 CN Benzamide, 2-[(3-(2-methoxyphenyl)-2-propynyl)methylamino]-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



RN 371258-57-0 CAPLUS
 CN Benzamide, 2-[(3-(4-methoxyphenyl)-2-propynyl)methylamino]-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 78 THERE ARE 78 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 23 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2001:246264 CAPLUS
 DOCUMENT NUMBER: 135:107296
 TITLE: Heteroannulation through copper catalysis: a novel and highly regio- and stereoselective cyclisation of alkynes leading to (E)-2-(2-arylvinyl)quinazolinones
 AUTHOR(S): Kundu, N. G.; Chaudhuri, G.
 CORPORATE SOURCE: Department of Organic Chemistry, Indian Association for the Cultivation of Science, Calcutta, Jadavpur, 700 032, India
 SOURCE: Tetrahedron Letters (2001), 42(15), 2883-2886
 CODEN: TELEAY; ISSN: 0040-4039
 PUBLISHER: Elsevier Science Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 135:107296
 ED Entered STN: 06 Apr 2001
 AB 2-(Alkylprop-2-ynylamino)benzamides reacted with aryl iodides under Pd-Cu catalysis to yield disubstituted alkynes, which underwent a novel cyclization in the presence of CuI, K₂CO₃, and Bu₄NBr in MeCN to yield (E)-1-alkyl-3-aryl-2-(2-arylvinyl)-4-quinazolinones in excellent yields

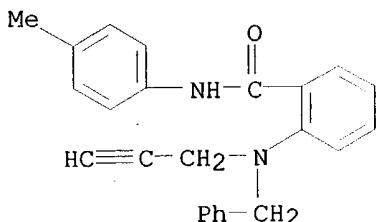
instead of the expected benzodiazepinones.

IT 350603-00-8

RL: RCT (Reactant); RACT (Reactant or reagent)
(prepn. of (arylvinyl)quinazolinones by regio- and stereoselective cyclization of (alkynylamino)benzamides)

RN 350603-00-8 CAPLUS

CN Benzamide, N-(4-methylphenyl)-2-[(phenylmethyl)-2-propynylamino]- (9CI)
(CA INDEX NAME)

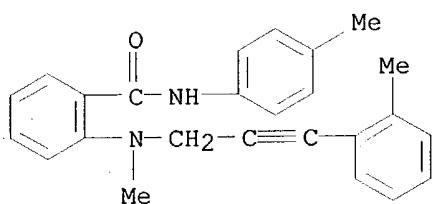


IT 350603-01-9P 350603-02-0P 350603-03-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of (arylvinyl)quinazolinones by regio- and stereoselective cyclization of (alkynylamino)benzamides)

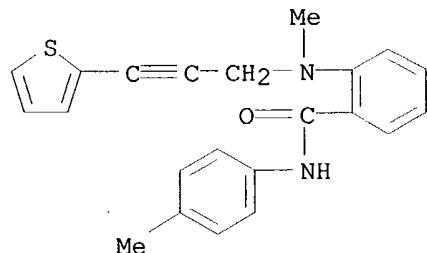
RN 350603-01-9 CAPLUS

CN Benzamide, 2-[(methyl[3-(2-methylphenyl)-2-propynyl]amino)-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



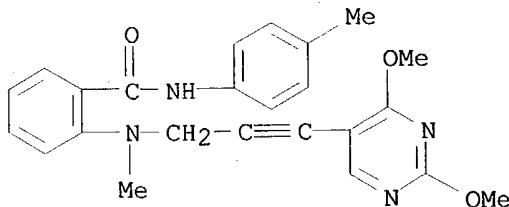
RN 350603-02-0 CAPLUS

CN Benzamide, N-(4-methylphenyl)-2-[(methyl[3-(2-thienyl)-2-propynyl]amino)- (9CI) (CA INDEX NAME)



RN 350603-03-1 CAPLUS

CN Benzamide, 2-[(3-(2,4-dimethoxy-5-pyrimidinyl)-2-propynyl)methylamino]-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 24 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2000:335388 CAPLUS

DOCUMENT NUMBER: 132:347491

TITLE: Preparation of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors

INVENTOR(S): Altmann, Karl-Heinz; Bold, Guido; Furet, Pascal; Manley, Paul William; Wood, Jeanette Marjorie; Ferrari, Stefano; Hofmann, Francesco; Mestan, Jurgen; Huth, Andreas; Kruger, Martin; Seidelmann, Dieter; Menrad, Andreas; Haberey, Martin; Thierauch, Karl-Heinz

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft m.b.H.; Schering Aktiengesellschaft

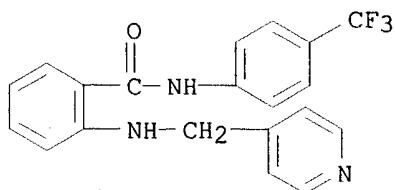
SOURCE: PCT Int. Appl., 77 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English

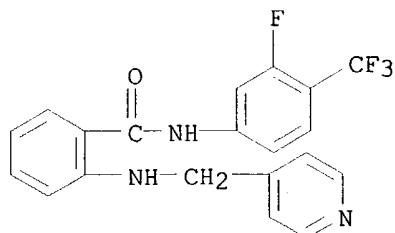
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

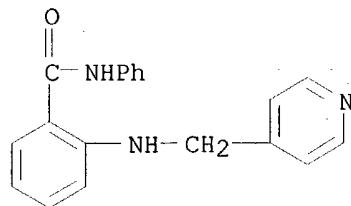
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000027820	A1	20000518	WO 1999-EP8545	19991108
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2346898	AA	20000518	CA 1999-2346898	19991108
BR 9915210	A	20010724	BR 1999-15210	19991108
EP 1129075	A1	20010905	EP 1999-971802	19991108
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2002529453	T2	20020910	JP 2000-581000	19991108
AU 758230	B2	20030320	AU 2000-13811	19991108
NZ 511339	A	20030725	NZ 1999-511339	19991108
NO 2001001894	A	20010704	NO 2001-1894	20010417
ZA 2001003290	A	20030123	ZA 2001-3290	20010423
US 2002019414	A1	20020214	US 2001-850434	20010507
US 6448277	B2	20020910		
ZA 2001004673	A	20020909	ZA 2001-4673	20010607
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PRIORITY APPLN. INFO.:			GB 1998-24579	A 19981110
			WO 1999-EP8545	W 19991108



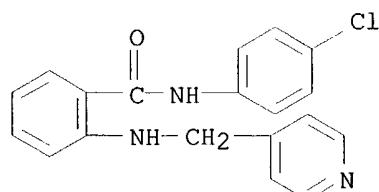
RN 269390-67-2 CAPLUS
CN Benzamide, N-[3-fluoro-4-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 269390-68-3 CAPLUS
CN Benzamide, N-phenyl-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 269390-69-4 CAPLUS
CN Benzamide, N-(4-chlorophenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



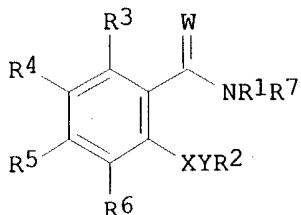
RN 269390-70-7 CAPLUS
CN Benzamide, N-(3-fluoro-4-methylphenyl)-2-[(4-pyridinylmethyl)amino]-, dihydrochloride (9CI) (CA INDEX NAME)

US 2001-850434 A3 20010507

OTHER SOURCE(S): MARPAT 132:347491

ED Entered STN: 19 May 2000

GI

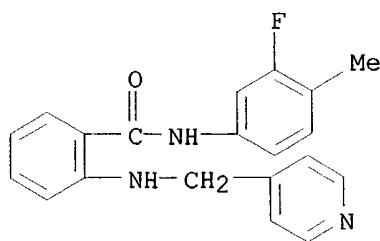


AB Use of title compds. I; W = O, S; X = NR8; Y = CR9R10(CH2)n, SO2; R9, R10 = H, alkyl; n = 0-3; R1 = aryl; R2 = mono- or bicyclic heteroaryl with the exception that R2 cannot = 2-phthalimidyl, and when Y = SO2 cannot represent 2,1,3-benzothiadiazol-4-yl; R3-R6 = H, substituent; R7, R8 = H, alkyl; or a N-oxide or a pharmaceutically acceptable salt thereof, for the prepn. of a pharmaceutical product for the treatment of a neoplastic disease which responds to an inhibition of the VEGF receptor tyrosine kinase activity is claimed. Thus, a mixt. of 4-pyridinecarboxaldehyde and 2-amino-N-(4-trifluoromethylphenyl)benzamide (prepn. given) in MeOH contg. HOAc was treated with NaBH3CN followed by 16 h stirring to give 2-[(4-pyridyl)methyl]amino-N-[4-(trifluoromethyl)phenyl]benzamide. Tested I inhibited Flt-1 VEGF receptor tyrosine kinase with IC50 = 0.18-0.56 .mu.M.

IT 269390-66-1P 269390-67-2P 269390-68-3P
 269390-69-4P 269390-70-7P 269390-71-8P
 269390-72-9P 269390-73-0P 269390-74-1P
 269390-75-2P 269390-76-3P 269390-77-4P
 269390-78-5P 269390-79-6P 269390-80-9P
 269390-81-0P 269390-82-1P 269390-83-2P
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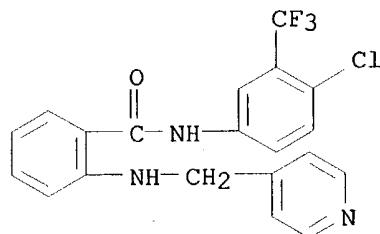
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors)

RN 269390-66-1 CAPLUS
 CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-[4-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

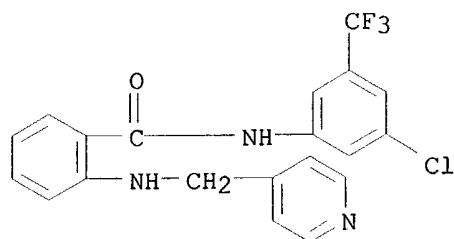


●2 HCl

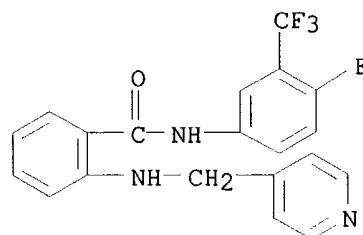
RN 269390-71-8 CAPLUS
CN Benzamide, N-[4-chloro-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 269390-72-9 CAPLUS
CN Benzamide, N-[3-chloro-5-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

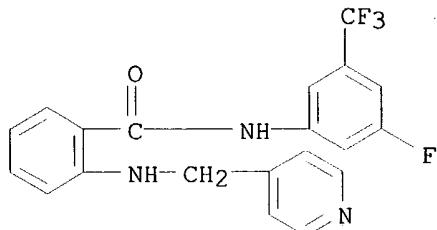


RN 269390-73-0 CAPLUS
CN Benzamide, N-[4-fluoro-3-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



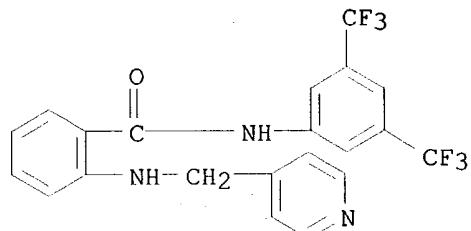
RN 269390-74-1 CAPLUS

CN Benzamide, N-[3-fluoro-5-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



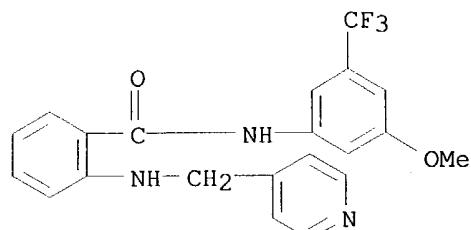
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CN Benzamide, N-[3,5-bis(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



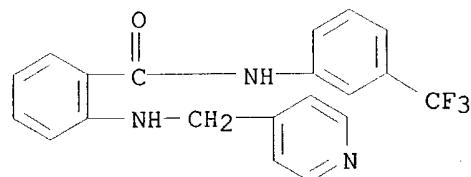
RN 269390-76-3 CAPLUS

CN Benzamide, N-[3-methoxy-5-(trifluoromethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



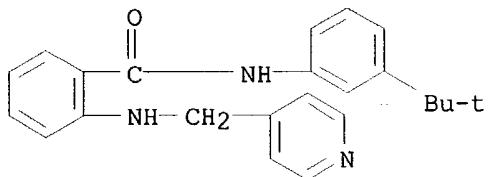
RN 269390-77-4 CAPLUS

CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



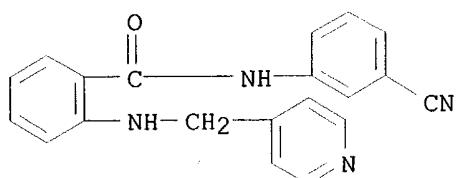
RN 269390-78-5 CAPLUS

CN Benzamide, N-[3-(1,1-dimethylethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



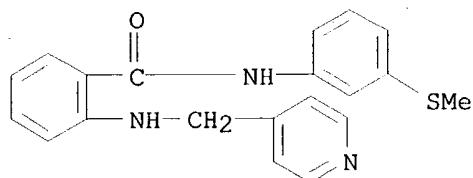
RN 269390-79-6 CAPLUS

CN Benzamide, N-(3-cyanophenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



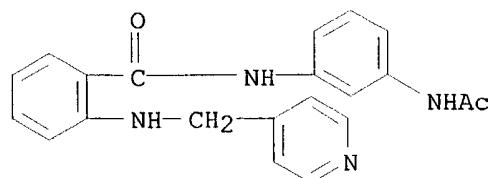
RN 269390-80-9 CAPLUS

CN Benzamide, N-[3-(methylthio)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



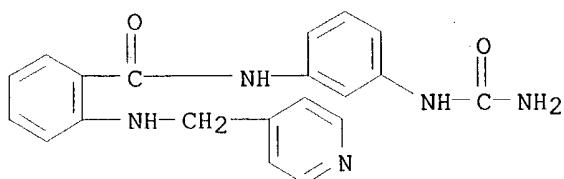
RN 269390-81-0 CAPLUS

CN Benzamide, N-[3-(acetylamino)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

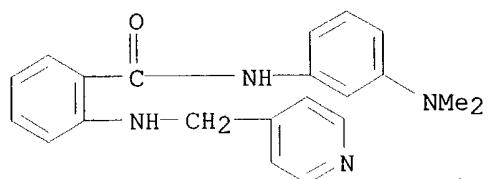


RN 269390-82-1 CAPLUS

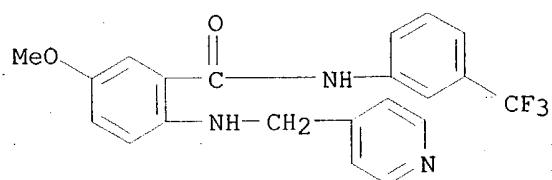
CN Benzamide, N-[3-[(aminocarbonyl)amino]phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



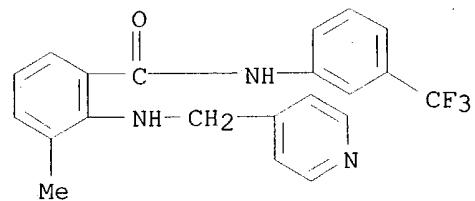
RN 269390-83-2 CAPLUS
 CN Benzamide, N-[3-(dimethylamino)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
 (CA INDEX NAME)



RN 269390-84-3 CAPLUS
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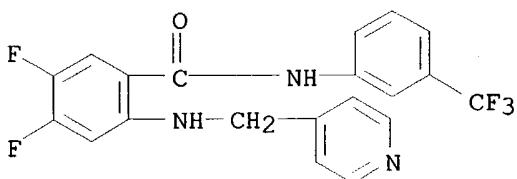


RN 269390-85-4 CAPLUS
 CN Benzamide, 3-methyl-2-[(4-pyridinylmethyl)amino]-N-[3-(trifluoromethyl)phenyl]-, dihydrochloride (9CI) (CA INDEX NAME)

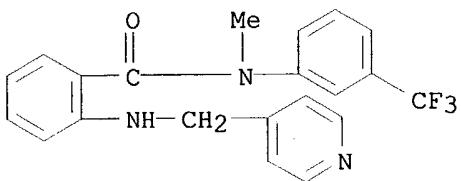


● 2 HCl

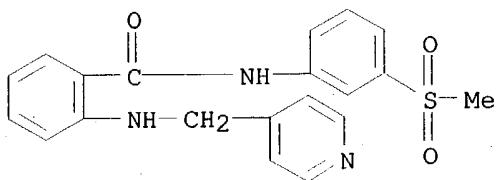
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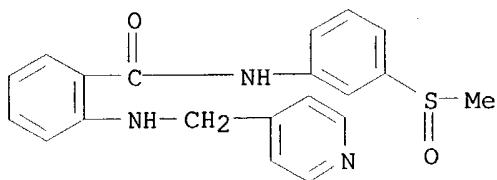
RN 269390-87-6 CAPLUS
CN Benzamide, N-methyl-2-[(4-pyridinylmethyl)amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



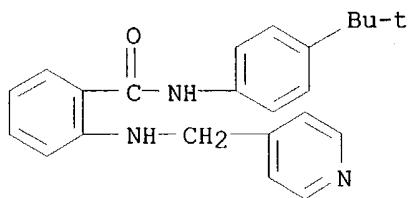
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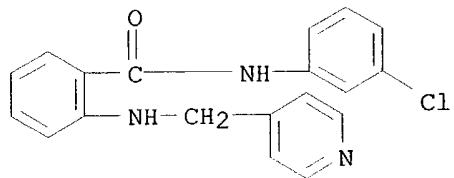
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CN Benzamide, N-[3-(methylsulfinyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



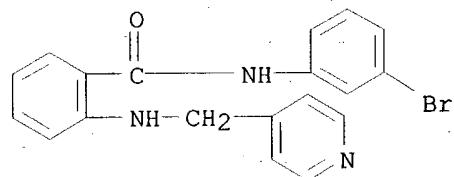
RN 269390-90-1 CAPLUS
CN Benzamide, N-[4-(1,1-dimethylethyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



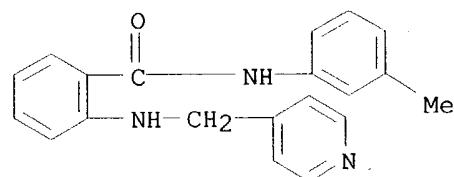
RN 269390-91-2 CAPLUS
CN Benzamide, N-(3-chlorophenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA
INDEX NAME)



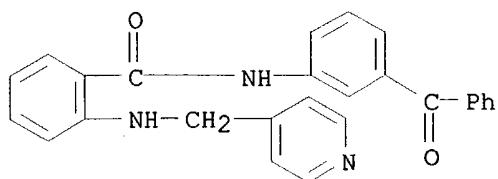
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INDEX NAME)



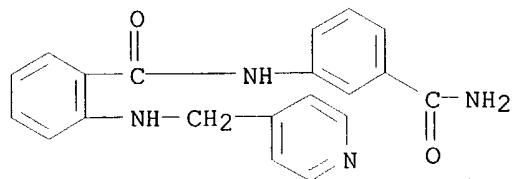
RN 269390-93-4 CAPLUS
CN Benzamide, N-(3-methylphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA
INDEX NAME)



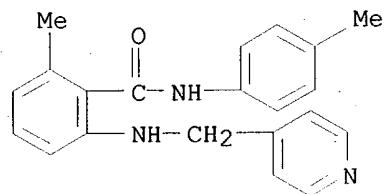
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CN Benzamide, N-(3-benzoylphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA
INDEX NAME)



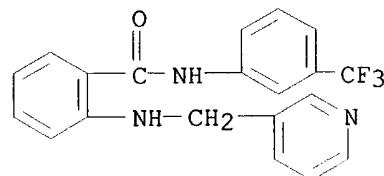
RN 269390-95-6 CAPLUS
CN Benzamide, N-[3-(aminocarbonyl)phenyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
(CA INDEX NAME)



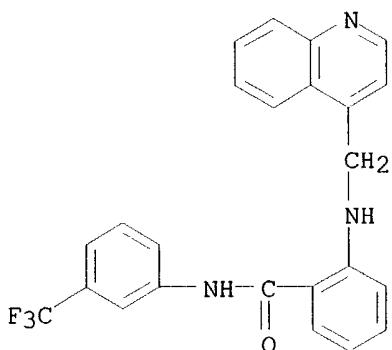
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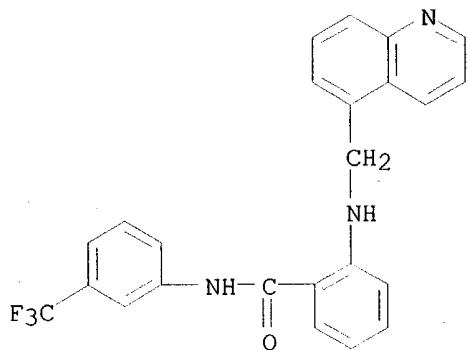
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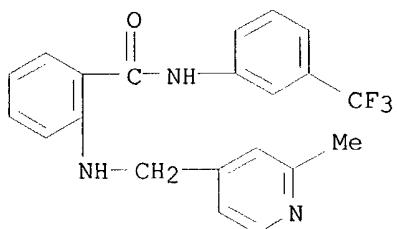
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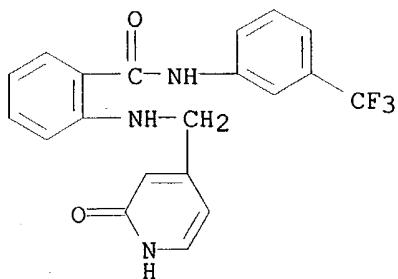
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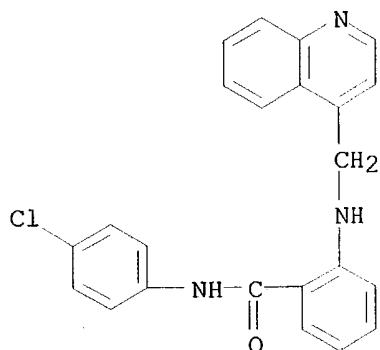
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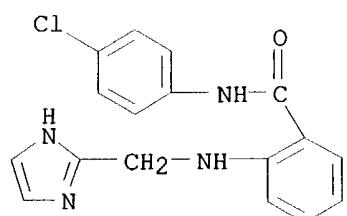
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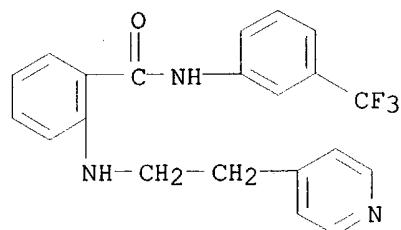
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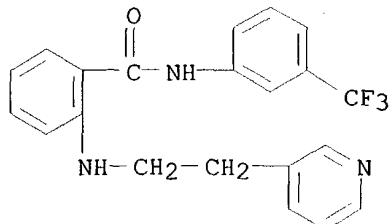
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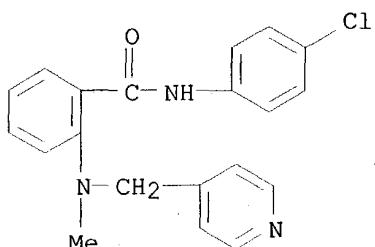
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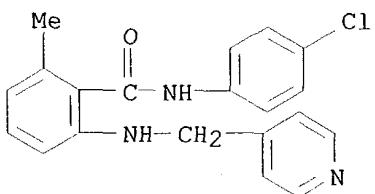
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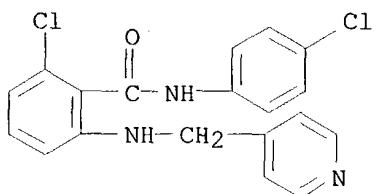
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(CA INDEX NAME)

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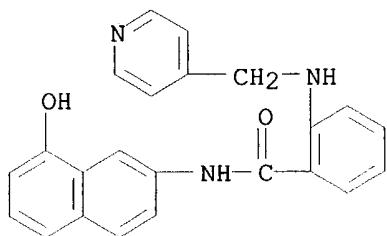
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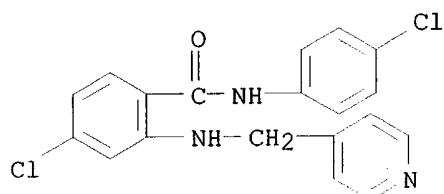
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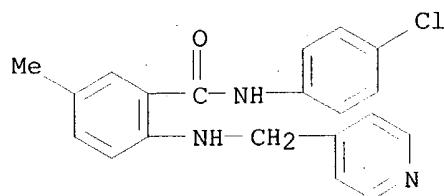
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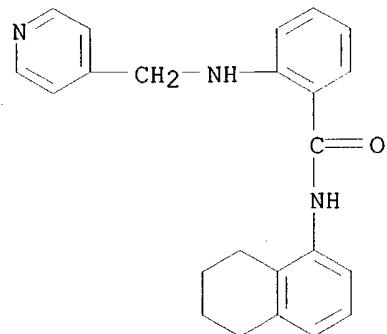
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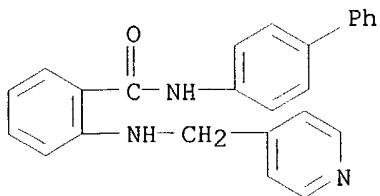
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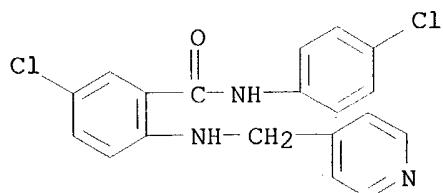
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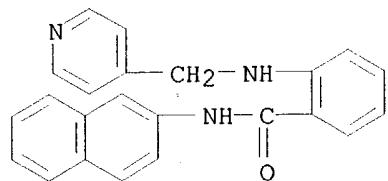
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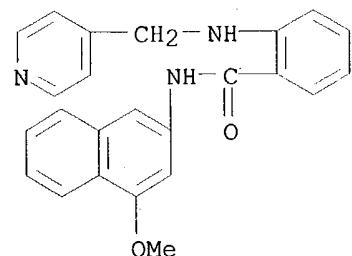
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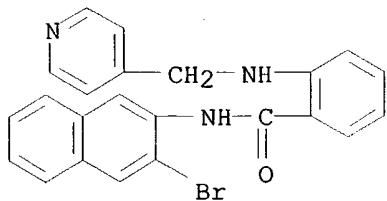
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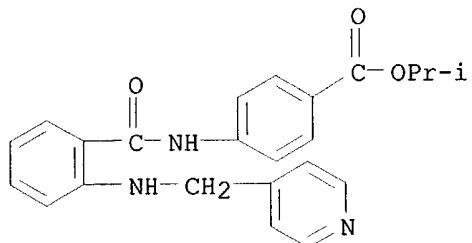


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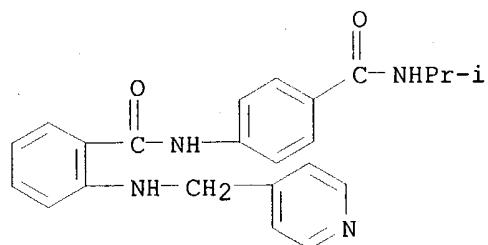
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CN Benzoic acid, 4-[(2-[(4-pyridinylmethyl)amino]benzoyl)amino]-, 1-methylethyl ester (9CI) (CA INDEX NAME)



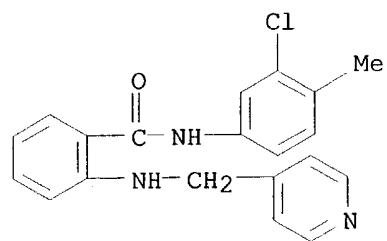
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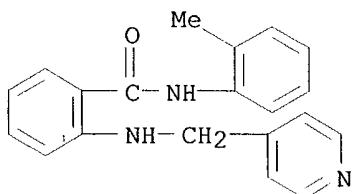
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CN Benzamide, N-(3-chloro-4-methylphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

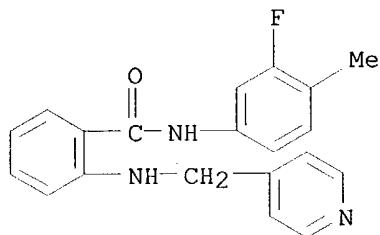


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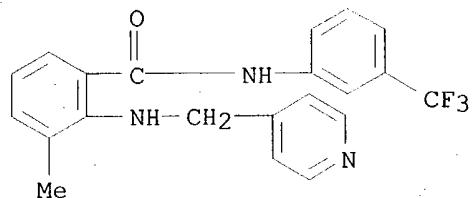
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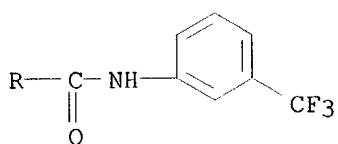
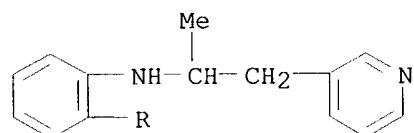
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 CN Benzamide, N-(3-fluoro-4-methylphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI)
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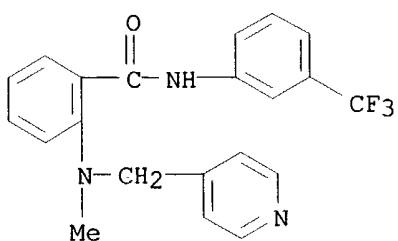
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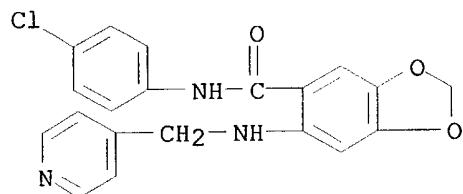
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 CN Benzamide, 2-[(1-methyl-2-(3-pyridinyl)ethyl)amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



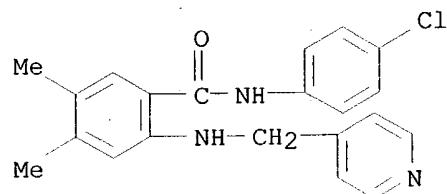
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 CN Benzamide, 2-[(1-methyl-2-(3-pyridinyl)ethyl)amino]-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)



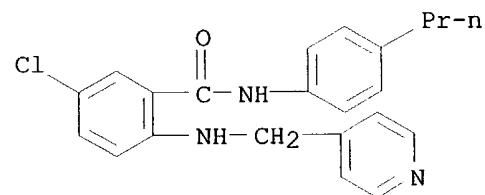
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CN 1,3-Benzodioxole-5-carboxamide, N-(4-chlorophenyl)-6-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



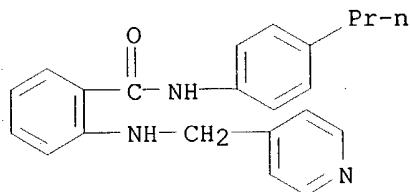
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CN Benzamide, N-(4-chlorophenyl)-4,5-dimethyl-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 269391-55-1 CAPLUS
CN Benzamide, 5-chloro-N-(4-propylphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

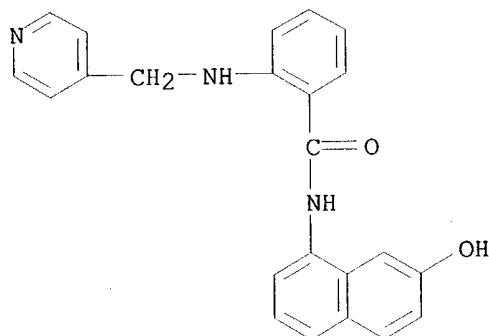


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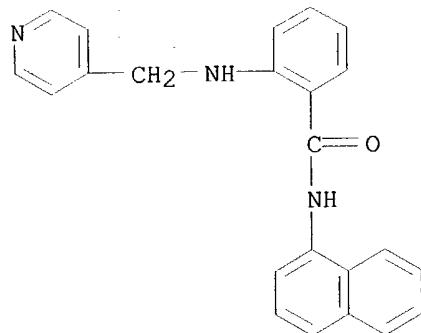
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CN Benzamide, N-(7-hydroxy-1-naphthalenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



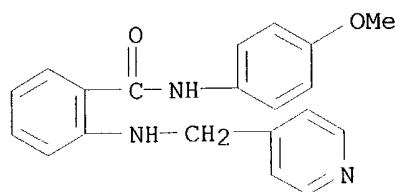
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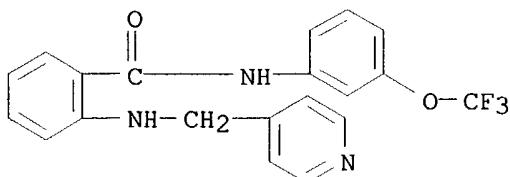
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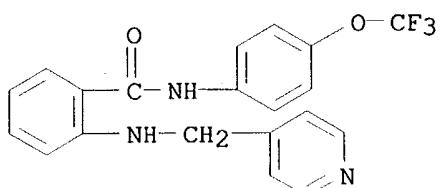
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CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-[3-(trifluoromethoxy)phenyl]-
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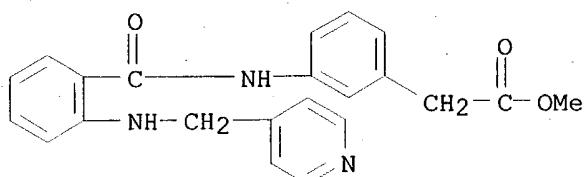
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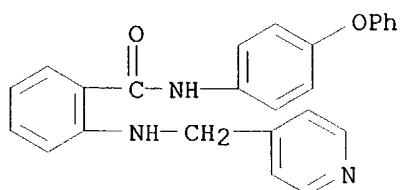
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CN Benzeneacetic acid, 3-[[2-[(4-pyridinylmethyl)amino]benzoyl]amino]-, methyl ester (9CI) (CA INDEX NAME)



RN 269391-63-1 CAPLUS

CN Benzamide, N-(4-phenoxyphenyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

14

THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L31 ANSWER 25 OF 55 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2000:335387 CAPLUS

DOCUMENT NUMBER: 132:334364

TITLE: Preparation of anthranilic acid amides as vascular endothelial growth factor receptor inhibitors.

INVENTOR(S):

Huth, Andreas; Seidelmann, Dieter; Thierauch, Karl-Heinz; Bold, Guido; Manley, Paul William; Furet, Pascal; Wood, Jeanette Marjorie; Mestan, Jurgen; Bruggen, Jose; Ferrari, Stefano; Kruger, Martin; Ottow, Eckhard; Menrad, Andreas; Schirner, Michael

PATENT ASSIGNEE(S):

Schering Aktiengesellschaft, Germany; Novartis Aktiengesellschaft

SOURCE:

PCT Int. Appl., 96 pp.

CODEN: PIXXD2

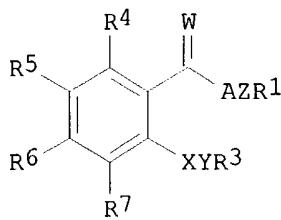
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

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WO 2000027819	A3	20000817		
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BR 9915553	A	20010814	BR 1999-15553	19991109
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EE 200100258	A	20021216	EE 2001-258	19991109
NO 2001002245	A	20010710	NO 2001-2245	20010507
BG 105588	A	20020430	BG 2001-105588	20010611
PRIORITY APPLN. INFO.:			GB 1998-24579 A	19981110
			DE 1999-19910396 A	19990303
			WO 1999-EP8478 W	19991109

OTHER SOURCE(S): MARPAT 132:334364

ED Entered STN: 19 May 2000

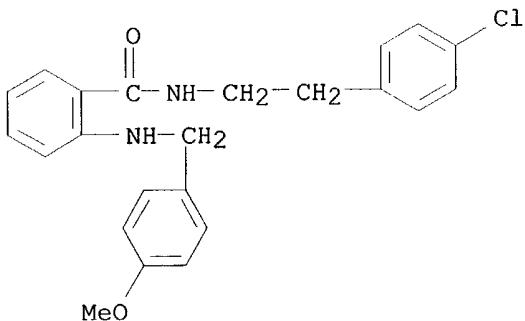
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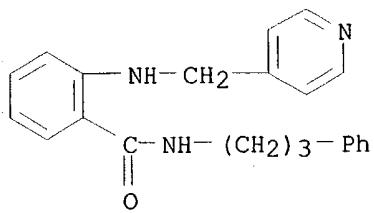
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N-(3-phenylprop-1-yl)-N2-(4-pyridylmethyl)anthranilamide. The latter inhibited VEGFR I with IC50 = 0.05 .mu.M.

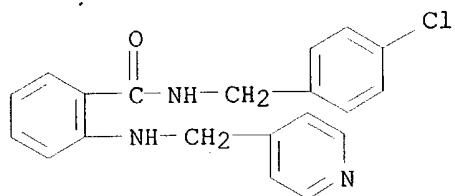
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 (prepn. of anthranilic acid amides as VEGF receptor inhibitors)
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 CN Benzamide, N-[2-(4-chlorophenyl)ethyl]-2-[(4-methoxyphenyl)methyl]amino]- (9CI) (CA INDEX NAME)



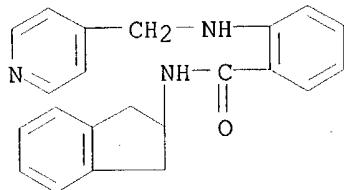
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 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of anthranilic acid amides as VEGF receptor inhibitors)
 RN 267891-04-3 CAPLUS
 CN Benzamide, N-(3-phenylpropyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



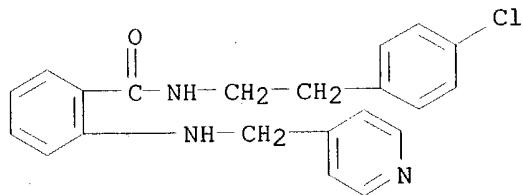
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CN Benzamide, N-[(4-chlorophenyl)methyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
(CA INDEX NAME)



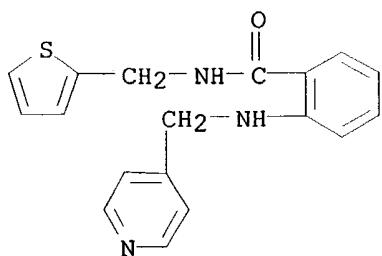
RN 267891-06-5 CAPLUS
CN Benzamide, N-(2,3-dihydro-1H-inden-2-yl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 267891-07-6 CAPLUS
CN Benzamide, N-[2-(4-chlorophenyl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
(CA INDEX NAME)

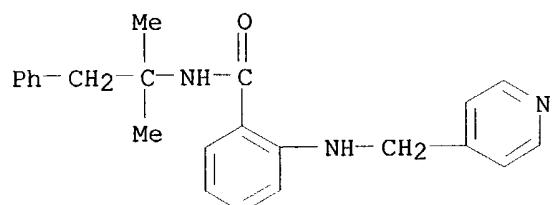


RN 267891-09-8 CAPLUS
CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-(2-thienylmethyl)- (9CI) (CA INDEX NAME)



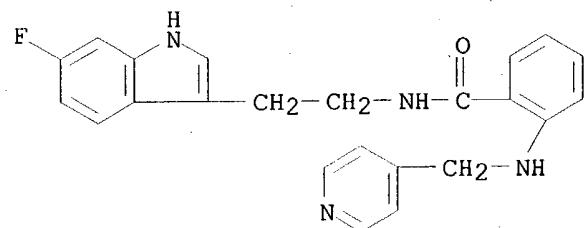
RN 267891-10-1 CAPLUS

CN Benzamide, N-(1,1-dimethyl-2-phenylethyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



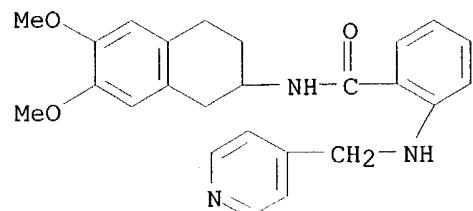
RN 267891-11-2 CAPLUS

CN Benzamide, N-[2-(6-fluoro-1H-indol-3-yl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



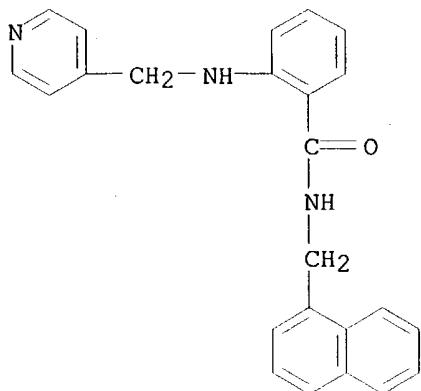
RN 267891-12-3 CAPLUS

CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-naphthalenyl)- (9CI) (CA INDEX NAME)



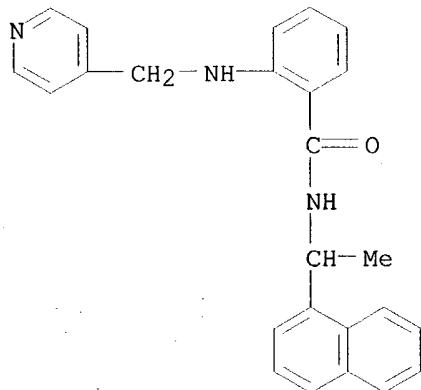
RN 267891-13-4 CAPLUS

CN Benzamide, N-(1-naphthalenylmethyl)-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



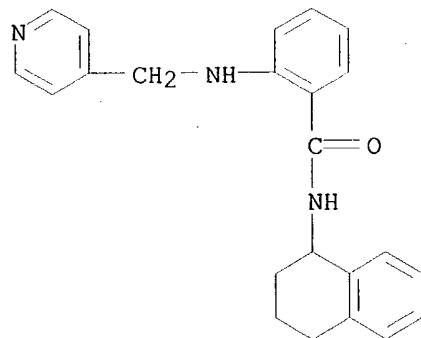
RN 267891-14-5 CAPLUS

CN Benzamide, N-[1-(1-naphthalenyl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
(CA INDEX NAME)



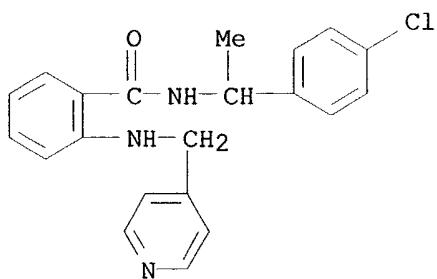
RN 267891-15-6 CAPLUS

CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-(1,2,3,4-tetrahydro-1-naphthalenyl)- (9CI) (CA INDEX NAME)

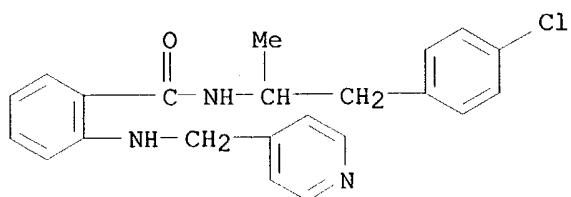


RN 267891-16-7 CAPLUS

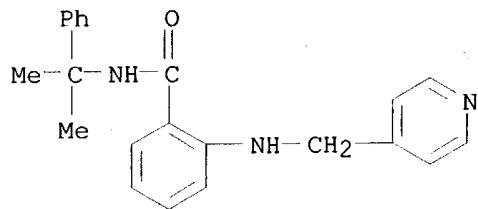
CN Benzamide, N-[1-(4-chlorophenyl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI)
(CA INDEX NAME)



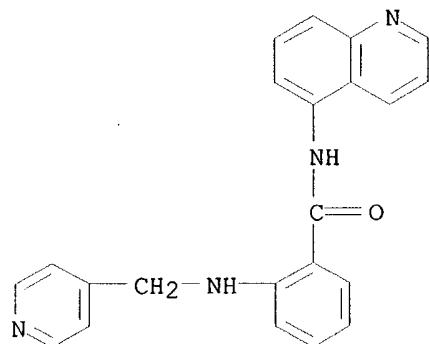
RN 267891-17-8 CAPLUS
 CN Benzamide, N-[2-(4-chlorophenyl)-1-methylethyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 267891-18-9 CAPLUS
 CN Benzamide, N-(1-methyl-1-phenylethyl)-2-[(4-pyridinylmethyl)amino]- (9CI)
 (CA INDEX NAME)

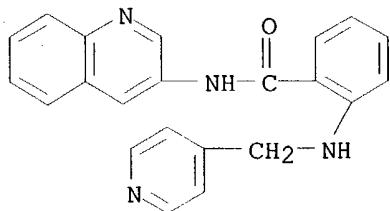


RN 267891-19-0 CAPLUS
 CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-5-quinolinyl- (9CI) (CA INDEX NAME)



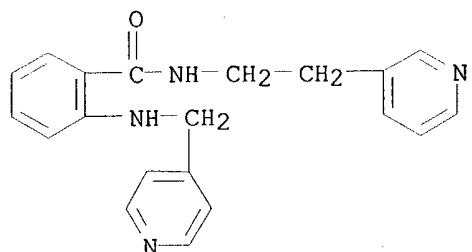
RN 267891-20-3 CAPLUS

CN Benzamide, 2-[(4-pyridinylmethyl)amino]-N-3-quinolinyl- (9CI) (CA INDEX NAME)



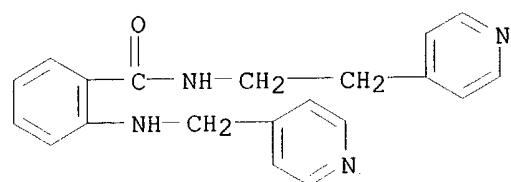
RN 267891-21-4 CAPLUS

CN Benzamide, N-[2-(3-pyridinyl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 267891-22-5 CAPLUS

CN Benzamide, N-[2-(4-pyridinyl)ethyl]-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)



RN 267891-23-6 CAPLUS

CN Benzamide, N-1-isoquinolinyl-2-[(4-pyridinylmethyl)amino]- (9CI) (CA INDEX NAME)

